



This is a digital copy of a book that was preserved for generations on library shelves before it was carefully scanned by Google as part of a project to make the world's books discoverable online.

It has survived long enough for the copyright to expire and the book to enter the public domain. A public domain book is one that was never subject to copyright or whose legal copyright term has expired. Whether a book is in the public domain may vary country to country. Public domain books are our gateways to the past, representing a wealth of history, culture and knowledge that's often difficult to discover.

Marks, notations and other marginalia present in the original volume will appear in this file - a reminder of this book's long journey from the publisher to a library and finally to you.

Usage guidelines

Google is proud to partner with libraries to digitize public domain materials and make them widely accessible. Public domain books belong to the public and we are merely their custodians. Nevertheless, this work is expensive, so in order to keep providing this resource, we have taken steps to prevent abuse by commercial parties, including placing technical restrictions on automated querying.

We also ask that you:

- + *Make non-commercial use of the files* We designed Google Book Search for use by individuals, and we request that you use these files for personal, non-commercial purposes.
- + *Refrain from automated querying* Do not send automated queries of any sort to Google's system: If you are conducting research on machine translation, optical character recognition or other areas where access to a large amount of text is helpful, please contact us. We encourage the use of public domain materials for these purposes and may be able to help.
- + *Maintain attribution* The Google "watermark" you see on each file is essential for informing people about this project and helping them find additional materials through Google Book Search. Please do not remove it.
- + *Keep it legal* Whatever your use, remember that you are responsible for ensuring that what you are doing is legal. Do not assume that just because we believe a book is in the public domain for users in the United States, that the work is also in the public domain for users in other countries. Whether a book is still in copyright varies from country to country, and we can't offer guidance on whether any specific use of any specific book is allowed. Please do not assume that a book's appearance in Google Book Search means it can be used in any manner anywhere in the world. Copyright infringement liability can be quite severe.

About Google Book Search

Google's mission is to organize the world's information and to make it universally accessible and useful. Google Book Search helps readers discover the world's books while helping authors and publishers reach new audiences. You can search through the full text of this book on the web at <http://books.google.com/>

UC-NRLF



QB 278 115

Chem.
General

LIBRARY

OF THE

University of California.

No.

3220

Division

Range

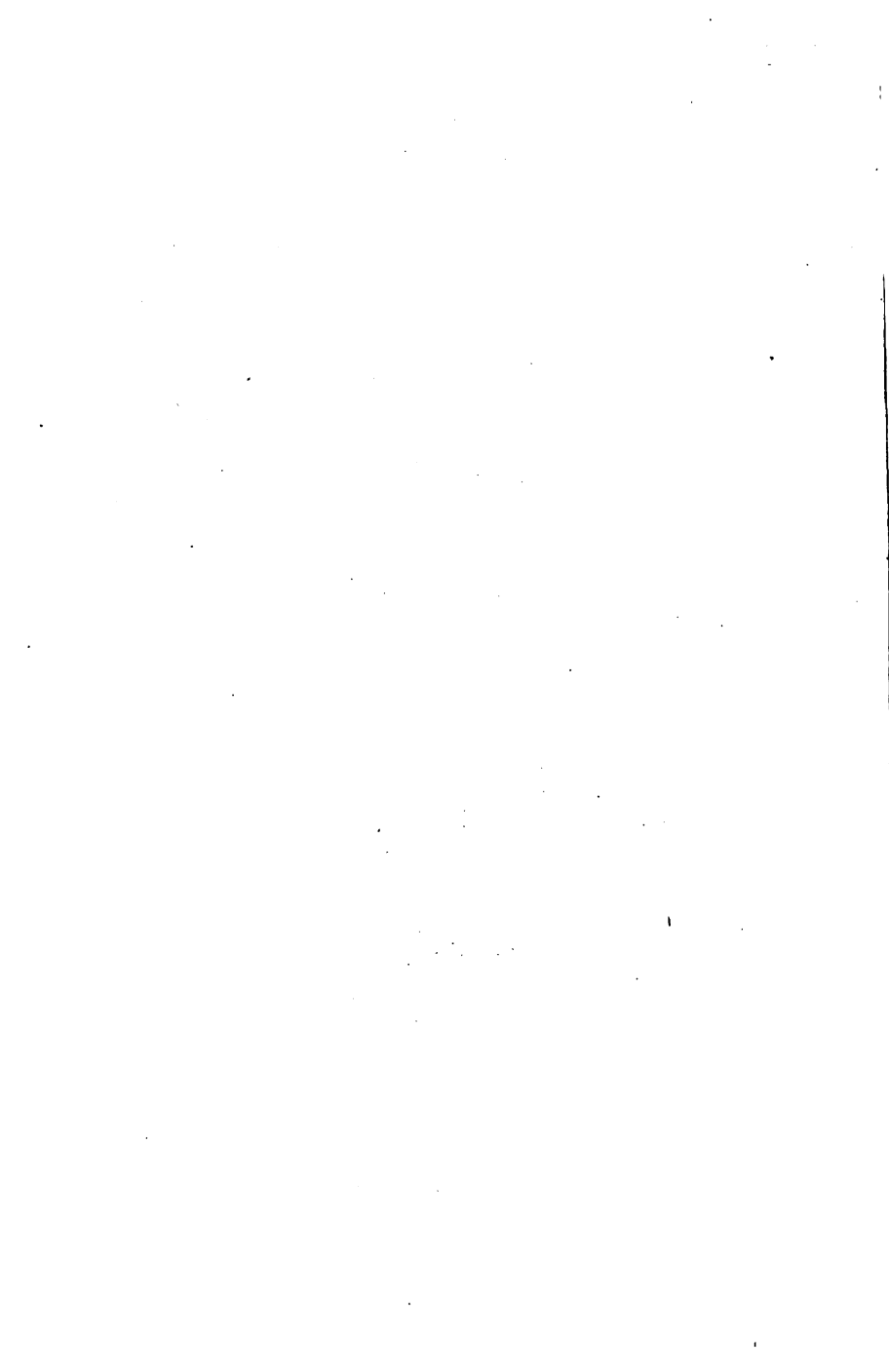
Shelf

Received

File

1870.





PRACTICAL CHEMISTRY.

LONDON: PRINTED BY
SPOTTISWOODE AND CO., NEW-STREET SQUARE
AND PARLIAMENT STREET

A COURSE
OF
PRACTICAL CHEMISTRY

ARRANGED FOR

THE USE OF MEDICAL STUDENTS.

BY

WILLIAM ODLING, M.B., F.R.S.

FELLOW OF THE ROYAL COLLEGE OF PHYSICIANS:

LECTURER ON CHEMISTRY AT ST. BARTHOLOMEW'S HOSPITAL.

FOURTH EDITION.

LONDON:
LONGMANS, GREEN, AND CO.
1869.

$\frac{Q.D=0}{04}$

PREFACE

TO

THE FOURTH EDITION.



IN this Fourth Edition of a Course of Practical Chemistry, arranged for the use of medical students, several minor improvements, suggested by further experience in teaching, have been effected.

The analytical portion of the work, in so far as regards the description of the methods employed, has been rearranged.

The new system of atomic weights and formulæ has been employed throughout; but it has not been thought necessary to introduce any general changes in the nomenclature.

W. O.

ST. BARTHOLOMEW'S HOSPITAL:

May, 1869.



PREFACE

TO

THE SECOND EDITION.



IN PREPARING a Second Edition of this Course of Practical Chemistry for the press, the first having been for upwards of three years out of print, many additions and alterations have been made, which it is believed will much increase its usefulness as a laboratory guide.

The first chapter, treating of chemical reactions and manipulation, is quite new; the second, relating to general analysis, has been rewritten; and the third and fourth chapters, treating respectively of toxicological and animal chemistry, have been carefully revised. In the second chapter more particularly, full explanations have been given of the tables for the examination of the several groups of bases and acids; so that the paragraphs relating to the individual members of the groups need scarcely be considered by the learner save for purposes of reference.

To maintain its adaptability to the wants of the medical student, the old scale of atomic weights has been exclusively employed throughout the body of the work.

In a short Appendix, however, some tables have been set up, in which the new atomic weights are used, with a view to illustrate the superior simplicity and mutual association of the modern unitary over the older dualistic formulæ. This Edition of the work is, moreover, illustrated by Mr. Branston with seventy woodcuts of microscopical preparations and chemical apparatus, all of them made expressly from drawings of the actual objects.

W. O.

ST. BARTHOLOMEW'S HOSPITAL:

June, 1865.

CONTENTS.

CHAPTER I.

INTRODUCTORY.

§ I.—CHEMICAL REACTIONS.

	PAGE
1. Equivalents	1
2. Atoms	2
3. Table of atomic weights . .	3
4. Symbolic notation	4
5. Acids and salts	6
6. Basicity of acids	7
7. Multivalent salts	8
8. Ammonium salts	9
9. Synoptic formulæ	10
10. Anhydrides	11
11. Double decomposition . .	12

§ II.—CHEMICAL MANIPULATION.

	PAGE
12. Sources of heat	13
13. Use of the blowpipe . . .	14
14. Glass-cutting and bending .	18
15. Glass-blowing and sealing .	19
16. Caoutchouc connectors &c. .	22
17. Construction of apparatus .	24
18. Filtration and decantation .	28
19. Supports and baths . . .	32
20. Heating liquids	35
21. Heating solids	39
22. Weighing and measuring . .	42
23. Specific gravities	47

CHAPTER II.

ANALYTICAL CHEMISTRY.

24. Course of operations . . .	52
--------------------------------	----

§ I.—BLOWPIPE EXAMINATION.

25. General effects	53
26. Special reactions	54

§ II.—SOLUTION AND PRECIPITATION.

27. Solution in water and acids .	61
28. Precipitation by reagents . .	62
29. Examination for basic groups	64

§ III.—EXAMINATION FOR BASES OF GROUP I.

30. Detection of first group . .	65
31. Table	68

32. Action of sulphuretted hydrogen	69
33. Sulphides soluble in sulphide of ammonium . .	70
34. Action of potash	71

§ IV.—EXAMINATION FOR BASES OF GROUP II.

35. Detection of second group . .	73
36. Table	76
37. Action of sulphide of ammonium	77
38. Action of potash	78
39. Action of ammonia	79
40. Behaviour of potash solution	81

§ V.—EXAMINATION FOR BASES OF GROUP III.

	PAGE
41. Detection of third group . . .	82
42. Table	83
43. Action of carbonate of ammonia	84
44. Bases unprecipitated by carbonate of ammonia	86

§ VI.—EXAMINATION FOR ACIDS.

45. General remarks	88
46. Table of preliminary results	89
47. Previous indications	90
48. Action of sulphuric acid	93
49. Application of liquid tests	94
50. Table	96
51. Action of nitrate of silver	96
52. Action of nitrate of barium	98
53. Action of chloride of calcium	98
54. Action of sulphate of magnesium	100
55. Action of ferric chloride	100
56. Order of testing for acids	101

§ VII.—SPECIAL SUBSTANCES.

57. General remarks	102
58. Insoluble compounds	102
59. Earthy salts	105
60. Oxides and sulphides	106
61. Acids or salts of hydrogen	107
62. Miscellaneous salts	108
63. Liquid or dissolved substances	109
64. Alkaline solutions of heavy metals	110

§ VIII.—INDIVIDUAL BASES OF GROUP I.

65. Tin	111
66. Arsenic	112
67. Antimony	113
68. Bismuth	113
69. Mercury	114
70. Lead	115

	PAGE
71. Silver	116
72. Copper	116
73. Cadmium	117

§ IX.—INDIVIDUAL BASES OF GROUP II.

74. Nickel	118
75. Cobalt	118
76. Manganese	119
77. Iron	120
78. Zinc	121
79. Chromium	122
80. Aluminium	122

§ X.—INDIVIDUAL BASES OF GROUP III.

81. Barium, strontium, calcium	123
82. Barium	123
83. Strontium	124
84. Calcium	124
85. Magnesium	125
86. Potassium	125
87. Ammonium	126
88. Sodium	126

§ XI.—REACTIONS OF INDIVIDUAL ACIDS.

89. Chromates	127
90. Nitrates	127
91. Chlorates	127
92. Carbonates	128
93. Sulphides and sulphydrates	129
94. Sulphites	130
95. Sulphates	130
96. Chlorides	130
97. Bromides	131
98. Iodides	131
99. Cyanides	132
100. Phosphates	132
101. Oxalates	133
102. Tartrates	134
103. Acetates	135
104. Benzoates	135
105. Borates	135
106. Fluorides	136
107. Silicates	137

CHAPTER III.

TOXICOLOGICAL CHEMISTRY.

	PAGE		PAGE
108. Introductory . . .	139	§ VII.—COPPER.	
§ I.—SULPHURIC ACID.		130. Dissolved . . .	155
109. Concentrated . . .	139	131. In organic mixtures . . .	156
110. Diluted . . .	140	§ VIII.—ARSENIC.	
111. In organic liquids . . .	141	132. Arsenious acid . . .	157
112. Stains on clothing . . .	141	133. Dissolved . . .	159
§ II.—NITRIC ACID.		134. Marsh's test . . .	161
113. Concentrated . . .	142	135. Original process . . .	162
114. Diluted . . .	142	136. Modified process . . .	164
115. Nitrates . . .	143	137. Reinsch's test . . .	165
116. In organic mixtures . . .	144	138. Impediments thereto . . .	167
§ III.—HYDROCHLORIC ACID.		139. Other forms of arsenic . . .	168
117. Concentrated . . .	144	140. In organic mixtures . . .	169
118. Diluted . . .	145	§ IX.—ANTIMONY.	
119. In organic mixtures . . .	146	141. Antimonial salts . . .	169
§ IV.—OXALIC ACID.		142. Dissolved . . .	170
120. Solid . . .	146	143. Marsh's test . . .	171
121. Dissolved . . .	147	144. Reinsch's test . . .	172
122. In organic mixtures . . .	147	145. In organic mixtures . . .	173
123. Insoluble salts . . .	149	§ X.—PRUSSIC ACID.	
§ V.—CORROSIVE SUBLIMATE.		146. Aqueous solution . . .	173
124. Solid . . .	149	147. In vapour . . .	174
125. Dissolved . . .	150	§ XI.—STRYCHNIA.	
126. In organic mixtures . . .	151	148. In pure state . . .	175
§ VI.—LEAD.		149. In organic mixtures . . .	178
127. Solid compounds . . .	153	§ XII.—MORPHIA.	
128. Dissolved . . .	154	150. In pure state . . .	179
129. In organic mixtures . . .	154	151. Opiate liquids . . .	180

CHAPTER IV.

ANIMAL CHEMISTRY.

§ I.—COMPOSITION OF TISSUES.		154. Ash of animal matter . . .	186
152. Organic and mineral constituents . . .	183	§ II.—NORMAL URINE.	
153. Ultimate organic constituents . . .	184	155. General properties . . .	188
		156. Urea . . .	189

	PAGE		PAGE
157. Uric acid . . .	100	168. Chemical testing . . .	206
158. Hippuric acid . . .	192	§ VI.—URINARY CALCULI.	
159. Colouring and extractive matter . . .	193	169. General characters . . .	207
160. Mineral salts . . .	194	170. Preliminary examination .	210
§ III.—ABNORMAL URINE.		171. Special tests . . .	211
161. Albuminous . . .	195	172. Systematic analysis . . .	213
162. Saccharine . . .	196	§ VII.—BLOOD.	
163. Biliary . . .	199	173. Coagulation . . .	215
164. Fatty . . .	199	174. Fibrin . . .	216
§ IV.—URINARY DEPOSITS.		175. Corpuseles . . .	217
165. Chemical . . .	201	176. Blood stains . . .	219
166. Organised . . .	203	177. Serum . . .	220
§ V.—CLINICAL EXAMINATION OF URINE.		§ VIII.—MISCELLANEOUS ANIMAL PRODUCTS.	
167. General examination . . .	205	178. Bile . . .	224
		179. Milk . . .	226
		180. Bone . . .	227

PRACTICAL CHEMISTRY.

CHAPTER I. INTRODUCTORY.

§ I.—CHEMICAL REACTIONS.

(1.) CHEMISTS are acquainted with about sixty different kinds of matter which have hitherto proved undecomposable, and are consequently termed simple bodies, or elements. These elements unite with one another in certain definite proportions to form an infinite variety of compounds; each particular chemical compound being always constituted of the same elements, combined together in the same proportion. Common salt, for instance, no matter how obtained, or when examined, is always found to consist of sodium and chlorine, united in the ratio of 23 parts by weight of the former element to 35·5 parts by weight of the latter.

The relative quantity of hydrogen which can enter into chemical combination being less than that of any other element, its combining proportion is taken as the standard of comparison or unity. It is found that 1 part by weight of hydrogen unites with 35·5 parts by weight of chlorine to form hydrochloric acid; and further that in very many chemical compounds 1 part of hydrogen may be displaced by 35·5 parts of chlorine to produce chlorine-derivatives of the respective compounds. Again, when hydrochloric acid is treated with metallic sodium, every 23 parts of sodium is found to expel 1 part of hydrogen and form common salt, by uniting with the 35·5 parts of chlorine previously combined with the 1 part of hydrogen.

The proportion of an element which unites with 1 part by weight of hydrogen, or which displaces 1 part by weight of hydrogen, to unite with 35·5 parts of chlorine, is called its equivalent. Thus 80 is the equivalent of bromine, 23 the equivalent

of sodium, and 39 the equivalent of potassium, because 80 parts of bromine, 23 parts of sodium, and 39 parts of potassium are respectively exchangeable for, or equivalent in combination to, 1 part of hydrogen.

(2.) But it is found in many cases that some multiple of the proportion of an element which unites with or displaces 1 part by weight of hydrogen, constitutes the smallest proportion of that element which actually enters into chemical combination. Thus although 25 parts of arsenic unite with 1 part of hydrogen to form arsenetted hydrogen, and with 35.5 parts of chlorine to form chloride of arsenic, yet it is allowed on all hands that the molecule* of arsenetted hydrogen contains three separable equivalents of hydrogen united with 75 parts of arsenic; and that the molecule of chloride of arsenic contains three separable equivalents of chlorine united with 75 parts of arsenic; and, in fact, that 75 parts of arsenic constitute the least indivisible proportion of arsenic which ever enters into a combination. The least indivisible proportion of an element which is found to enter into chemical combination is termed its atom, and the number expressing that proportion is called its atomic weight. Hence the atomic weight of an element sometimes coincides with its equivalent weight, as in the case of sodium, and is sometimes a multiple of its equivalent weight, as in the case of arsenic.

* The determination of the molecule of a compound body is based upon very many considerations, physical as well as chemical, which cannot be fully entered into here. The most important of these considerations relate to specific heat, atomic volume, direct combination, analogy, mode of derivation, and, above all, to metamorphoses by substitution. Thus the molecules of marsh gas, ammonia, water, and hydrochloric acid, might each be represented with one equivalent of hydrogen. But the molecule of marsh gas is represented with four equivalents of hydrogen, because in it $\frac{1}{4}$ or $\frac{2}{4}$ or $\frac{3}{4}$ or $\frac{4}{4}$ of the hydrogen can be displaced by substitution at four successive stages. The molecule of ammonia is represented with three equivalents of hydrogen, because in it $\frac{1}{3}$ or $\frac{2}{3}$ or $\frac{3}{3}$ of the hydrogen may be displaced at three successive stages. The molecule of water is represented with two equivalents of hydrogen, because in it $\frac{1}{2}$ or $\frac{2}{2}$ of the hydrogen can be displaced at two successive stages; while the molecule of hydrochloric acid is represented with one equivalent of hydrogen, because in it the hydrogen must be displaced at once or not at all.

The determination of equivalents is a purely experimental question, which, in the majority of instances, has been answered with almost absolute exactitude; but the determination of atomic weights is a question of judgment, to which in many cases very different answers were until lately accorded. Chemists are now agreed, however, as to the atomic weights of all the most important elements, and their agreement extends equally to atomic weights which are multiples of, as to those which are identical with, the equivalents of the respective elements. It is only with regard to a few of the less known elements that any great difference of opinion now exists as to the correlations of their respective equivalents and atomic weights.

(3.) The following tables exhibit lists of the most important elements, with their accepted atomic weights, their symbols or abbreviated names, and the names and symbols of their principal compounds with hydrogen or chlorine. All the hydrides are volatile, and when in the gaseous state occupy the same volume.

Element			Hydride	
Atomic weight	Symbol	Name	Symbol	Name
1	H	Hydrogen	H ₂	
19	F	Fluorine	HF	Hydrofluoric acid
35.5	Cl	Chlorine	HCl	Hydrochloric acid
80	Br	Bromine	HBr	Hydrobromic acid
127	I	Iodine	HI	Hydriodic acid
16	O	Oxygen	H ₂ O	Water
32	S	Sulphur	H ₂ S	Sulphuretted hydrogen
14	N	Nitrogen	H ₃ N	Ammonia
31	P	Phosphorus	H ₃ P	Phosphoretted hydrogen
11	B	Boron	H ₃ B	Hydride of boron?
12	C	Carbon	H ₄ C	Marsh-gas
28	Si	Silicon	H ₄ Si	Silicated hydrogen

Perissad Metal, &c.			Artiad Metal, &c.		
Atomic weight	Name	Chloride	Atomic weight	Name	Chloride
18	Ammonium	NH_4Cl	40	Calcium	Ca Cl_2
1	Hydrogen	H Cl	87.5	Strontium	Sr Cl_2
7	Lithium	Li Cl	137	Barium	Ba Cl_2
23	Sodium	Na Cl	24	Magnesium	Mg Cl_2
39	Potassium	K Cl	65	Zinc	Zn Cl_2
108	Silver	Ag Cl	112	Cadmium	Cd Cl_2
200	Mercury	Hg Cl^*	200	Mercury	Hg Cl_2
63.5	Copper	Cu Cl^*	63.5	Copper	Cu Cl_2
27.5	Aluminum	Al Cl_3^*	59	Nickel	Ni Cl_2
52.5	Chromium	Cr Cl_3^*	59	Cobalt	Co Cl_2
56	Iron	Fe Cl_3^*	56	Iron	Fe Cl_2
75	Arsenic	As Cl_3	55	Manganese	Mn Cl_2
122	Antimony	Sb Cl_3	118	Tin	$\text{Sn Cl}_2 \text{ \& } \text{Sn Cl}_4$
210	Bismuth	Bi Cl_3	207	Lead	Pb Cl_2
196	Gold	$\left\{ \begin{array}{l} \text{Au Cl}_3 \\ \text{Au Cl} \end{array} \right.$	197	Platinum	$\text{Pt Cl}_2 \text{ \& } \text{Pt Cl}_4$

It is observable that the metals mercury, copper, iron, tin, gold, and platinum, form two distinct chlorides, and, as will be afterwards seen, two distinct sets of oxides and salts, corresponding respectively thereto. Several of the other metals also form two chlorides and two sets of salts, but only the chlorides shown in the table, and their corresponding salts, are familiarly known. Cuprous, aurous, and platinous salts, moreover, are for the most part very unstable, and rarely met with save when specially prepared.

(4.) Each symbol, as exemplified in the above tables, represents one combining proportion, or atom, of the element. Thus N stands not for nitrogen in general, but for 14 parts by weight of

* It is doubtful whether the molecules and consequent formulæ of some or all of the chlorides corresponding to the formulæ marked with asterisks should not be doubled. The words 'perissad' and 'artiad' are applied to both metals and non-metals, accordingly as they combine with an odd or even number of atoms of chlorine or hydrogen.

nitrogen as compared with 1 part by weight of hydrogen. Chemical combination is represented by the apposition of symbols; thus KCl signifies a compound of 39 parts of potassium and 35.5 parts of chlorine united to form 74.5 parts of chloride of potassium. A small figure placed to the right of a symbol indicates a multiple quantity of that particular element. Thus HgCl_2 , or corrosive sublimate, signifies a compound of one atom of mercury with two atoms of chlorine. A larger figure placed to the left of an allocation of symbols multiplies the entire compound. Thus 3HNO_3 represents three proportions of nitric acid, a compound body consisting of one atom of hydrogen, one atom of nitrogen, and three atoms of oxygen. The single formula of a compound body represents the atom, or smallest indivisible proportion, of that particular compound, and its atomic weight is the sum of the atomic weights of its constituents. Thus the atomic weight of nitric acid $\text{HNO}_3 = 1 + 14 + 16 \times 3 = 63$.

In writing the formulæ of compound bodies, it is sometimes found advisable to break them up in different ways—by means of periods, brackets, parentheses, &c. Thus for sulphuric acid we sometimes write $\text{H}_2\text{O}.\text{SO}_3$, instead of H_2SO_4 ; for nitrate of ammonium, $\text{NH}_4.\text{NO}_3$, instead of $\text{N}_2\text{H}_4\text{O}_3$; and for phosphate of calcium $\text{Ca}_3(\text{PO}_4)_2$, instead of $\text{Ca}_3\text{P}_2\text{O}_8$. Sometimes these breaks are purely arbitrary or conventional, and should then be dispensed with as much as possible. At other times they indicate a real molecular isolation of one portion of a compound from the remainder, when their use is perfectly legitimate.

The symbol for the atom of an element is sometimes marked with one or more dashes, to indicate its equivalency or interchangeable value for hydrogen. For instance, Ag is sometimes marked with a single dash to show that the atom of silver may be substituted for an atom of hydrogen, so as to combine with an atom of chlorine, thus $\text{Ag}'\text{Cl}$. Again, Pb is marked with two dashes, and Bi with three dashes to indicate that the atoms of lead and bismuth may be respectively substituted for two and three atoms of hydrogen, so as to combine with two and three atoms of chlorine, thus $\text{Pb}''\text{Cl}_2$ and $\text{Bi}'''\text{Cl}_3$.

The signs +, -, and = are used almost in their ordinary algebraical sense. The sign + signifies addition to, or rather mixture with; the sign - subtraction from; and the sign = equivalency with, or rather conversion into. Thus the equation $2\text{HCl} + \text{CuO} = \text{CuCl}_2 + \text{H}_2\text{O}$, implies that an atom of hydrochloric acid, mixed with an atom of oxide of copper, yields an atom of chloride of copper together with an atom of water. Or the equation may of course be written $2\text{HCl} + \text{CuO} - \text{H}_2\text{O} = \text{CuCl}_2$. In modern chemistry the sign + is no longer used to express combination.

(5.) Those hydrogenised compounds which can readily exchange some or all of their hydrogen for its equivalent of metal constitute the acids. From habit one particular reaction is adopted as the conventional criterion of acidity, namely that effected by the hydrates of potassium and sodium. An acid is simply a hydrogenised body which, when treated with hydrate of potassium, can exchange hydrogen for potassium with simultaneous formation of water, thus:



The solutions of such bodies are generally found to have the power of reddening blue litmus paper, and of effervescing with alkaline carbonates, so that the possession of these properties may be looked upon as more or less characteristic of an acid. Oxygenised acids, such as the nitric HNO_3 , and acetic $\text{H}_4\text{C}_2\text{O}_4$, are also called ternary, while non-oxygenised acids, such as the hydrochloric HCl , and sulphydric, H_2S , are called binary. These binary acids were formerly known as hydracids. In most ternary acids the number of oxygen atoms is two, or three, or four, as shown below:

HClO_2 Chlorous.	HClO_3 Chloric.	HClO_4 Perchloric.
HNO_2 Nitrous.	HNO_3 Nitric.	HI O_4 Periodic.
HBO_2 Boracic.	$\text{H}_2\text{S O}_3$ Sulphurous.	$\text{H}_2\text{S O}_4$ Sulphuric.
$\text{H}_4\text{C}_2\text{O}_4$ Acetic	$\text{H}_2\text{C O}_3$ Carbonic.	$\text{H}_2\text{C}_2\text{O}_4$ Oxalic.
$\text{H}_3\text{P O}_4$ Hypophos.	$\text{H}_3\text{P O}_3$ Phosphorous.	$\text{H}_3\text{P O}_4$ Phosphoric.

It is not uncommon to have one or more of the hydrogen atoms

of a ternary acid which are not replaceable by metal exchanged for chlorine, and some or all of its oxygen atoms exchanged for sulphur. Thus we have acetic acid $\text{H}_4\text{C}_2\text{O}_4$, chloracetic acid $\text{HCl}_3\text{C}_2\text{O}_4$, and sulphacetic acid $\text{H}_4\text{C}_2\text{S}_2$; with their corresponding salts, $\text{NaH}_3\text{C}_2\text{O}_4$ acetate, $\text{NaCl}_3\text{C}_2\text{O}_4$ chloracetate, and $\text{NaH}_3\text{C}_2\text{S}_2$ sulphacetate of sodium, for instance.

(6.) Acids in which only one atom of hydrogen can be replaced by metal are called monobasic. The principal monobasic acids with which the student will have to deal are the following :

H Cl	Hydrochloric.	Na Cl	Chloride of sodium.
H Cl O_3	Chloric.	Na Cl O_3	Chlorate of sodium.
H Br	Hydrobromic.	Na Br	Bromide of sodium.
H I	Hydro-iodic.	Na I	Iodide of sodium.
H N O_3	Nitric.	Na N O_3	Nitrate of sodium.
H B O_2	Boracic.*	Na B O_2	Borate of sodium.
$\text{H}_4\text{C}_2\text{O}_4$	Acetic.	$\text{NaH}_3\text{C}_2\text{O}_4$	Acetate of sodium.

Acids in which two atoms of hydrogen can be replaced by metal are called dibasic. The most important of them are comprised in the following table :

H ₂ O	Hydric acid	.	.	{	KHO K ₂ O	Potash. Oxide of potassium.
H ₂ S	Sulphydric acid	.		{	NaHS Na ₂ S	Sulphydrate of sodium. Sulphide of sodium.
H ₂ SO ₃	Sulphurous acid	.		{	NaHSO ₃ Na ₂ SO ₃	Acid sulphite of sodium. Sulphite of sodium.
H ₂ SO ₄	Sulphuric acid	.		{	KHSO ₄ K ₂ SO ₄	Acid sulphate of K. Sulphate of potassium.
H ₂ SiO ₃	Silicic acid	.	.	.	K ₂ SiO ₃	Silicate of potassium.
H ₂ CO ₃	Carbonic acid	.		{	KHCO ₃ K ₂ CO ₃	Acid carbonate of K. Carbonate of potassium.
H ₂ C ₂ O ₄	Oxalic acid	.	.	{	KHC ₂ O ₄ K ₂ C ₂ O ₄	Acid oxalate of K. Oxalate of potassium.
H ₂ C ₄ O ₆	Tartaric acid	.	.	{	KH ₅ C ₄ O ₆ KNaH ₄ C ₄ O ₆	Cream of tartar. Rochelle salt.

* Another variety of boracic acid has the formula H_3BO_3 , and is tribasic.

Until lately many of these acids were considered as monobasic and represented by the halves of the formulæ here given; but the evidence of their dibasicity is at present indisputable. This class of bodies includes sulphydric acid, or sulphuretted hydrogen, the sulphur analogue of water, which itself often plays the part of an acid, and is included in the foregoing list.

Acids in which three atoms of hydrogen may be replaced by metal are called tribasic, the most important of which are—

H_3PO_4	Phosphoric acid	.	$\left\{ \begin{array}{l} KH_2PO_4 \\ Na_2HPO_4 \\ Ag_3PO_4 \end{array} \right.$	Phosphate of potassium. Phosphate of sodium. Phosphate of silver.
H_3AsO_4	Arsenic acid	.	Ag_3AsO_4	Arseniate of silver.
$H_3C_6O_7$	Citric acid	.	$Ag_3H_3C_6O_7$	Citrate of silver.

Monometallic and dimetallic arseniates and citrates are also familiarly known.

The monometallic salts of dibasic and tribasic acids closely resemble the acids themselves in their action on blue litmus paper and on alkaline hydrates and carbonates. They constitute, indeed, a mere variety of the class of acids.

In polyhydrogenised acids, it does not follow that the units of basicity are equal to the units of hydrogen,—acetic acid $H_4C_2O_4$, tartaric acid $H_6C_4O_6$, and citric acid $H_8C_6O_7$, for example, being but monobasic, dibasic, and tribasic respectively, or capable of exchanging respectively but one, two, and three atoms of hydrogen for metal.

(7.) In the illustrative salts above adduced, each atom of hydrogen in the acid has been displaced by one atom of univalent metal. Thus we had nitrate of potassium KNO_3 , derived from nitric acid HNO_3 ; oxalate of sodium $Na_2C_2O_4$, derived from oxalic acid $H_2C_2O_4$; phosphate of silver Ag_3PO_4 , derived from phosphoric acid H_3PO_4 , &c. &c. But the salts of multivalent metals have usually, though not invariably, a somewhat greater complexity of constitution. Their chlorides, derived from two or three atoms of hydrochloric acid, are given on page 4, and their other salts derived from monobasic acids are found to correspond closely with their chlorides. Thus dichloride and dini-

trate of tin, $\text{Sn}''\text{Cl}_2$ and $\text{Sn}''(\text{NO}_3)_2$, are derived from two molecules of hydrochloric and nitric acid, H_2Cl_2 and $\text{H}_2(\text{NO}_3)_2$, respectively; trichloride and trinitrate of bismuth, $\text{Bi}'''\text{Cl}_3$ and $\text{Bi}'''(\text{NO}_3)_3$, derived from three molecules of hydrochloric and nitric acid, H_3Cl_3 and $\text{H}_3(\text{NO}_3)_3$, respectively. But the formulæ of the salts of dibasic acids with divalent metals and of tribasic acids with triquivalent metals are very simple. Thus lead sulphate $\text{Pb}''\text{SO}_4$ is derived from sulphuric acid H_2SO_4 , and bismuth phosphate $\text{Bi}'''\text{PO}_4$, from phosphoric acid H_3PO_4 , &c. On the other hand the salts of dibasic acids with triquivalent metals, and of tribasic acids with divalent metals, are highly complex. Sulphate of bismuth, for example, must be represented by the formula $\text{Bi}'''_2(\text{SO}_4)_3$, derived from three atoms of sulphuric acid $\text{H}_6(\text{SO}_4)_3$, and so in other instances.

(8.) It will be perceived from the above tables and examples, that a salt is usually derived from its corresponding acid by a substitution of metal for hydrogen. Many salts, however, known as salts of the alkaloids, are formed in a different way—namely, by a direct union of the acid with ammonia or some other alkaloidal base; as exemplified by hydrochloride of ammonia NH_3HCl , nitrate of ammonia NH_3HNO_3 , &c. But salts of this character may also be considered to contain a composite metal, or rather metalloid, in place of the hydrogen of the acid. Thus by associating with each atom of ammonia in the salt an atom of hydrogen from the acid, each such atom of ammonia NH_3 , becomes converted into an atom of ammonium NH_4 , a pseudo-metallic grouping, which in its combinations presents a most marked analogy to potassium, as illustrated below:

Ammonia salts.	Ammonium salts.	Potassium salts.
NH_3HCl Hydrochlor.	NH_4Cl Chloride.	KCl Chloride.
NH_3HNO_3 Nitrate.	NH_4NO_3 Nitrate.	KNO_3 Nitrate.
$\text{NH}_3\text{H}_2\text{SO}_4$ Acid sulph.	NH_4HSO_4 Acid sulph.	KHSO_4 Acid sulph.
$(\text{NH}_3)_2\text{H}_2\text{SO}_4$ Sulphate.	$(\text{NH}_4)_2\text{SO}_4$ Sulphate.	K_2SO_4 Sulphate.

Without assuming any knowledge of the actual molecular arrangement of ammoniacal salts, it is found most convenient in

practice, especially when comparing them with metallic salts, to regard them as salts of ammonium rather than as salts of ammonia. But the salts of the correlated alkaloids aniline, morphia, strychnia, &c., are usually represented after the manner of ammonia salts, thus :

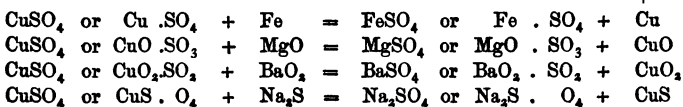
Hydrochlorides.			Acetates.		
NH_3	$\cdot \text{HCl}$	Ammonia.	NH_3	$\cdot \text{H}_4\text{C}_2\text{O}_2$	Ammonia.
$\text{C}_6\text{H}_7\text{N}$	$\cdot \text{HCl}$	Aniline.	$\text{C}_6\text{H}_7\text{N}$	$\cdot \text{H}_4\text{C}_2\text{O}_2$	Aniline.
$\text{C}_{17}\text{H}_{19}\text{NO}_3$	$\cdot \text{HCl}$	Morphia.	$\text{C}_{17}\text{H}_{19}\text{NO}_3$	$\cdot \text{H}_4\text{C}_2\text{O}_2$	Morphia.
$\text{C}_{21}\text{H}_{23}\text{N}_2\text{O}_2$	$\cdot \text{HCl}$	Strychnia.	$\text{C}_{21}\text{H}_{23}\text{N}_2\text{O}_2$	$\cdot \text{H}_4\text{C}_2\text{O}_2$	Strychnia.

(9.) It was formerly the custom to regard ternary acids as compounds of one or more equivalents of water with a special oxidised grouping, and the corresponding salts as compounds of one or more equivalents of metallic oxide with the same oxidised grouping. Thus the formulæ of chloric acid, and chlorate of potassium were written $\text{H}_2\text{O}.\text{Cl}_2\text{O}_3$ and $\text{K}_2\text{O}.\text{Cl}_2\text{O}_3$, corresponding to 2HClO_3 and 2KClO_3 , respectively. But this custom, which was based on assumptions since shown to be erroneous, is now falling gradually into disuse. Thus the acids of chlorine form the following series, the ternary members of which may be obtained by direct and successive oxidation of the binary member, hydrochloric acid, the hydrogen of which cannot possibly exist in the state of water :

HCl	Hydrochloric.	KCl	Chloride.
HClO	Hypochlorous.	KClO	Hypochlorite.
HClO_2	Chlorous.	KClO_2	Chlorite.
HClO_3	Chloric.	KClO_3	Chlorate.
HClO_4	Perchloric.	KClO_4	Perchlorate.

It is true that many ternary acids and salts may be directly or indirectly formed from, or separated into, a special oxidised grouping and water or metallic oxide, but the same acids and salts may also be formed from, or separated into, a variety of other sub-compounds, and the one mode of composition or decomposition has no more right than has each of the others to set up for itself a rational formula. Thus if we act upon sulphate of

copper by metallic iron, magnesia, peroxide of barium, and sulphide of sodium respectively, we have the following reactions :



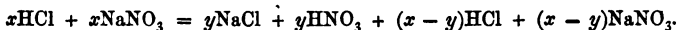
From each of these reactions there might, with equal reason, be inferred the pre-existence in sulphate of copper of the respective groupings SO_4 , SO_3 , SO_2 , and CuS ,—and the correctness of the respective rational formulæ $\text{Cu} \cdot \text{SO}_4$, $\text{CuO} \cdot \text{SO}_3$, $\text{CuO}_2 \cdot \text{SO}_2$, and $\text{CuS} \cdot \text{O}_4$, in accordance with the theories of Dulong, Berzelius, Longchamps, and Laurent respectively. Moreover, sulphate of copper may be electrolysed into Cu and $\text{SO}_3 + \text{O}$, while it may be formed by combining CuO with SO_3 , or CuO_2 with SO_2 , or CuS with O_4 . From considerations of this kind chemists have thought it better to employ, as much as possible, what are called synoptic formulæ, which express only the composition of acids and salts, and not their internal molecular arrangement.

(10.) The anhydrous acids assumed to pre-exist in ternary acids and salts are mostly hypothetical, but those which have an actual independent existence are found to be quite devoid of acid properties, and in their reactions upon various classes of bodies to differ greatly from the corresponding acids. Hence the appellation acid is altogether inapplicable to them, and consequently the phrase anhydrous acid is become gradually disused, and the word anhydride adopted in its stead. The only anhydrides often concerned in chemical reactions are the carbonic, silicic, stannic, sulphurous, and arsenious. With the carbonic and sulphurous anhydrides may be associated carbonic oxide and sulphuric anhydride respectively, as in the following table :

CO	Carbonic oxide.
CO_2	Carbonic anhydride.
SiO_2	Silica.
SnO_2	Stannic anhydride.
SO_2	Sulphurous anhydride.
SO_3	Sulphuric anhydride.
As_2O_3	Arsenious anhydride.

The carbonic, silicic, stannic, sulphurous, and arsenious acids are unstable ill-defined bodies, which readily break up into water and the respective anhydrides.

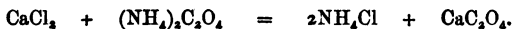
(11.) The general term *salt* is often taken to include the acid, or salt of hydrogen, as well as the salt of a true metal, such as sodium, or of a quasi-metal, such as ammonium. Using the term in this broad sense, it may be said that whenever different salts of different bases or basylides occur in solution, they undergo mutual decomposition to a greater or less extent according to circumstances. Thus when solutions of chloride of hydrogen and nitrate of sodium are mixed together in equivalent proportions, we have produced some chloride of sodium and nitrate of hydrogen together with some unaltered chloride of hydrogen and nitrate of sodium, or the two salts become four salts, thus:



But whenever any one of the freshly formed salts is removed from the sphere of chemical action by precipitation or volatilisation, there is a complete instead of a partial decomposition, thus:



Hence we arrive at the following general law. Any two salts which, by an exchange of their respective basylides, can, under the conditions of the experiment, form an insoluble or volatile compound, undergo a complete double decomposition with precipitation of the insoluble or evolution of the volatile compound. Oxalate of calcium, for instance, being insoluble in water, we know that when chloride of calcium solution is mixed with excess of oxalate of ammonium solution, the whole of the calcium will be precipitated in the form of oxalate of calcium, thus:



Again, acetic acid or acetate of hydrogen being volatile at a moderate temperature, we know that when acetate of sodium is

heated with sulphate of hydrogen a double decomposition will take place, and acetic acid be liberated, thus :



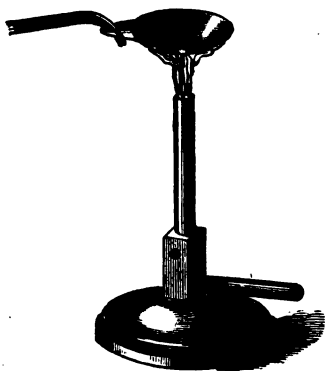
As will hereafter be seen, the deposition of characteristic precipitates and evolution of characteristic gases or vapours constitute the most general means by which the presence of particular bodies can be analytically established.

§ II.—CHEMICAL MANIPULATION.

(12.) The spirit-lamp is very useful for many operations, especially when a small, smokeless, not over hot flame is required. The charcoal burner is scarcely necessary in a laboratory furnished with gas, but is otherwise almost indispensable. In a chauffer of the kind figured on page 38, one or two pieces of charcoal may be kept slowly burning, by occasionally blowing off the ash; or a large brisk fire may be kept up, capable of boiling a gallon or more of water. A few pieces of charcoal may also be readily burnt on a coarse wire grating or trellis, resting by its edges on a couple of bricks or other suitable support. But gas is by far the most convenient fuel for ordinary laboratory work. Among burners which are very generally useful may be mentioned the bat's-wing-gauze burner. This consists of a large bat's-wing nipple, screwed into an elbow of brass tube standing on a flat iron foot, and provided with a gallery of some kind to support a brass chimney covered at the top with wire gauze. The bat's-wing flame burnt without the chimney is convenient for bending glass tube, and when reduced by partially turning off the gas, is well fitted for blowpipe testing (fig. 4). With the chimney on, and the ascending mixture of gas and air burnt at the top of the gauze, as shown on page 34, a large smokeless flame is obtained suitable for heating sand-baths, flasks, test-tubes, &c. An argand burner, screwed into a flat iron foot and provided with a short brass chimney, is also convenient for

many purposes, especially when a steady long-continued heat is required. When small flasks, &c., have to be heated, a flat brass ring with an aperture about the size of a shilling may be placed on the top of the argand chimney so as to confine the heat. In the Bunsen burner (fig. 1), gas issues from a short jet fixed in the interior of an upright tube, having holes at the bottom through which air is sucked in, so as to produce a mixture of gas and air to be burnt at the top of the tube. The flame is

Fig. 1.



smokeless, and from its great heat well suited for ignitions on a small scale. In many Bunsen burners the gas may be either burnt in a single upright flame, as above described, or in a flat rosette of smaller jets well adapted for evaporations, distillations, &c., as shown on page 37. The Bunsen burner may be further modified by slipping into its upright tube, a somewhat narrower tube, hav-

ing an expanded, broad, flat opening,—the flat flame from which is especially suitable for blowpipe use. Herapath's burner (fig. 2), is a very useful instrument for effecting strong ignitions or fusions, and for glass working. It consists of a large blowpipe jet *a*, connected with the mouthpiece by a flexible tube, and sliding in the interior of a brass tube *b*, furnished with a supply of gas, through a lateral projection *c*, fitting on to an elbow on which it moves easily, so as to allow of the flame being turned in any direction.

(13.) The mode of using the mouth blowpipe, though difficult to describe, is fortunately very easy to learn. The best blowpipe for general laboratory use is that designed by Dr. Black (fig. 3 *a*).

It consists of a small conical tube of brass or tin plate closed at its broad, and open at its narrow end, which is usually furnished with a mouthpiece of bone or ivory. From the side of the cone

Fig. 2.



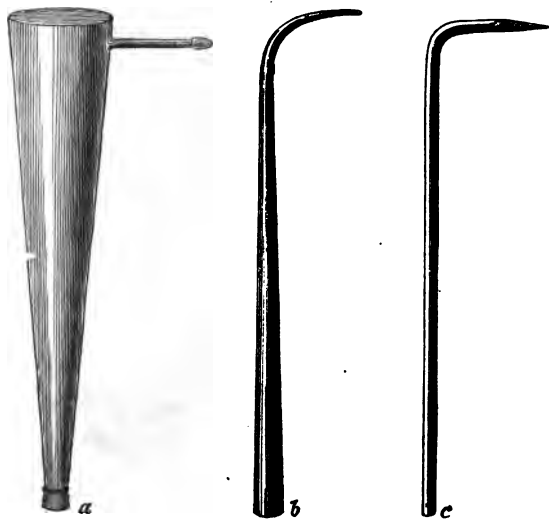
near its broad end, there projects a piece or narrow tube about an inch long terminating in a jet, through which the current of air issues. This jet or nozzle should be turned out of a solid piece of metal, should be strictly conical both inside and out, and should fit on to the conical end of the narrow tube by mere pressure without screwing. It is advisable to have a fine jet for testing, and a coarse jet for glass working and general heating; but for these last purposes a common brazier's blowpipe (fig. 3 *b*), or even a bent glass tube drawn out to a fine point (fig. 3 *c*) will usually suffice.

A coarse bat's-wing burner with the gas partly cut off, so as to produce a flame scarcely larger than that of a candle, will be found very convenient for blowpipe testing; but in default of gas, the flame of a large candle, or of a spirit lamp fed with solution of turpentine in spirits of wine, may be employed. The candle-

wick should be cut of medium length, and turned in the same direction as the blowpipe jet.

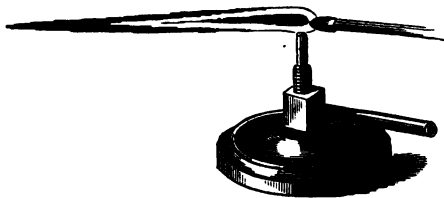
In using a blowpipe, the air must be projected by the muscles, not of the chest, but of the mouth, which should be blown

Fig. 3.



out like a trumpeter's. The breathing has to be carried on through the nose only; and, after a little practice, it will be found easy to keep up a continuous pressure with the cheeks and lips

Fig. 4.

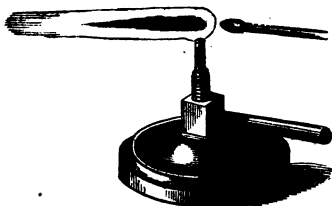


quite unaffected by the alternating respiratory movements of the trunk. The mouth should never be allowed to get exhausted of air, but be constantly reple-

nished from behind through the nose. It will be requisite for the student to acquire the power of producing either an oxidising or reducing flame at will. To produce an oxidising flame (fig. 4),

the extremity of the jet should be placed inside the gas- or candle-flame, and a steady not too forcible current of air maintained. By this means a continuous, silent, lateral flame, of sharply defined conical form, will be produced. The interior of this flame close to the jet will be dark, and contain an excess of cold air from the mouth; while surrounding it will be seen a somewhat thick layer of clear blue. This is the cone of perfect combustion, the pointed extremity of which constitutes the hottest part of the flame. Outside this blue cone, more especially at its termination, will

Fig. 5.



be perceived a very slightly luminous yellow cone, in which the external air is in excess and at a very high temperature. This constitutes the oxidising portion of the flame, by exposure to which, a small piece of metallic tin, the size of a pin's head, should gradually swell up into a pulverulent mass of peroxide of tin. To produce a reducing or deoxidising flame (fig. 5), the blowpipe jet, which should be rather fine, is placed just outside the gas- or candle-flame, and a somewhat forcible current of air maintained. The resulting blowpipe flame is much less sharply defined than that previously described, and consists principally of a large luminous cone containing an excess of unconsumed carbon, which exerts a powerful reducing action. A minute portion of peroxide of tin, heated in this flame on a charcoal support, may be readily brought to the metallic state even without the use of a flux.

The heating power of the blowpipe flame depends upon the continuous impulsion of hot gaseous matter on to the substance under examination, and upon its rapid removal so soon as it has imparted its heat by contact. Moreover, a very perfect combustion of the fuel is effected, partly by air projected through the jet, partly by external air drawn in from the sides of the jet and flame, and coinciding in direction with the projected stream.

(14.) Some amount of skill must be attained by the chemical student in constructing apparatus of glass tubing, and in otherwise working with glass. Ordinary tubing or rod may be cut of any required length by making a firm scratch across it with a triangular file, and then breaking it sharply at the file-mark by a conjoint pull and bend. Glass may also be, as it were, sawn through by means of a file, but the operation is rather tedious. It may be much facilitated by occasionally wetting the file with turpentine, or even with water. Glass may be bored through in a similar manner by a drill or a hard bradawl dipped in turpentine. When the neck of a flask, rim of a beaker, or portion of other glass apparatus, becomes cracked, a piece of ignited charcoal, or preferably of pastille, held in contact with the glass immediately in front of the crack, will serve to extend it in any desired direction, so as to cut off the neck or rim, &c. The charcoal may be kept alight by gently blowing on it. Pastilles are made of charcoal powder formed into a mass with thick gum, and rolled into quill-sized sticks, which are dusted with charcoal and dried.

Glass stirring rods are best made from a piece of solid rod or cane long enough for two stirrers. This is to be heated at its middle point in the blowpipe flame, turning it constantly round and gently extending it, until a constriction is formed, which when cold is scratched with a file. The two halves are then snapped apart, and the ends of the stirrers rounded off by directing the flame upon them for a few moments.

Subliming or reduction tubes are made by taking a piece of glass tubing of about 0.2 inch bore and six inches long, heating it at its middle point in the blowpipe flame until the glass is thoroughly softened, and then gently pulling the two halves asunder. The glass of which these tubes are formed should be clear, thin, and difficultly fusible.

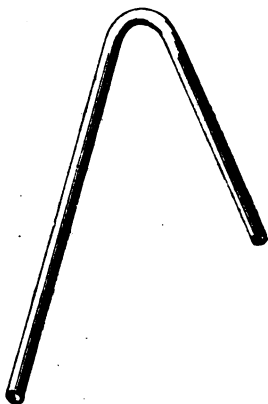
Ordinary glass tubing, unless very thick, may be bent in the flame of a bat's-wing or Bunsen burner. The piece of tubing should be heated over a considerable portion of its length, either at once or successively, and as it gradually softens be bent into

the required shape. The production of any constriction in the bore may be avoided by causing some length of tube to take part in the bend, which should never form a sharp angle, but a well-rounded curve, as in the syphon (fig. 6).

Glass tubing may be drawn out to an almost capillary point, as in the jet of a wash bottle, end of a pipette, &c., by heating a portion of it, about a quarter of an inch long, until it just softens and then steadily pulling until a sufficient constriction is produced, which is afterwards scratched with a file and snapped across. The point may have its edges rounded, and its aperture further diminished if necessary, by holding it in the flame for a few seconds or so. In order to point the extremity of a piece of tubing, another piece of tubing or rod must first be joined to it by the blowpipe, so that the necessary extension may be made. It is often advisable to thicken a tube slightly at the spot where it is to be drawn out, by rotating it for some time in the flame, and gently pressing its ends together. In this way the conical aperture may be made both strong and fine.

(15.) Glass-blowing is a valuable accomplishment to the practical chemist, but there are only one or two small operations with which it is actually necessary for the student to become familiar. It will be rarely worth while for him to make his own test-tubes, but he should be able to reseal any that have got accidentally broken. By means of the blowpipe, a piece of waste rod or tubing must first be joined firmly on to two or three projecting points of the broken end, and be made to coincide as nearly as practicable with the axis of the test-tube, as in fig. 7.

Fig. 6.



Then, at about half an inch or more from its broken end, the tube is to be steadily heated in a large blowpipe-flame, and constantly rotated until a considerable constriction is formed, when gentle extension may be employed. The flame is next to be directed upon what will form the bottom of the tube, just at its

Fig. 7.



bend or shoulder, and the extension continued. By this means the end to be pulled away will be left irregularly conical, and that of the new tube well rounded. Finally, the thread of glass proceeding from the cone is to be strongly heated at its junction with the tube until it melts and becomes detached. There is thus always left on the bottom of the tube a little burr or projection of melted glass, which, if sufficiently small, may be made to disappear by heating it and the whole bottom of the tube until the glass is soft, and then blowing into the tube with moderate force while rotating it in the mouth. The burr, if too large for this treatment, may be melted in a small blowpipe-flame, and then have a piece of waste tubing, previously warmed, applied to it, and quickly drawn away, so as to bring the burr with it. In this manner a smaller burr will be left, which may be melted into the bottom of the tube as above described. The mouth of a test-tube may be everted by softening it in the flame, and then bending the edge uniformly outwards with the smooth end of an old file previously heated nearly to redness; or else a conical piece of charcoal may be twisted into the softened mouth of the tube.

Glass tubes sealed at both ends constitute the best means of preserving small specimens, either of liquid or solid. One end of the tube is first sealed as if for a test-tube, save that the small burr need not be interfered with. The other end is then constricted to a greater or less extent, according to the size of the specimen to be introduced, so as to leave a funnel-shaped

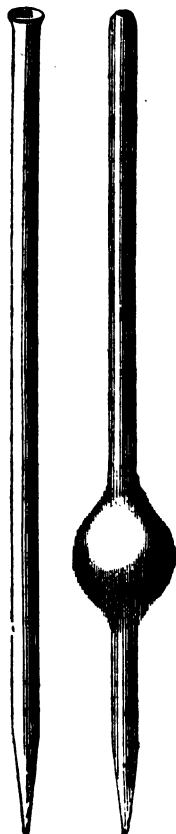
appendage, as in fig. 8. Liquids may be introduced through a very constricted opening by alternately warming and cooling the body of the tube while the liquid is contained in the funnel. After the substance has been introduced, a small blowpipe-flame is gradually brought to bear upon the constriction, and the sealing completed. Very careful heating is more particularly required when the constriction is moist from the passage of liquid. For practice the student should seal up small tubes filled to within an inch or so of their length with water, spirit, sulphur, sugar, &c. When sealed tubes are used for effecting reactions under pressure, their ends have to be made with great care so as not to be less resisting than the original sides.

When a small bulb has to be blown upon the end of a piece of tubing, the closed end must first be thickened by rotating it in the flame for some time, and pressing it up with a piece of charcoal until enough glass has accumulated. This thick portion has next to be strongly heated, then withdrawn from the flame, and have air quickly but gently blown into it from the other end, during constant rotation of the tube in the mouth. Or a bulb may be blown in the course of a tube, as in making a bulbed pipette (fig. 9 *b*), for instance. For this purpose a portion of the tube is to be thickened considerably by rotating it in the

Fig. 8.



Fig. 9.



flame for some time and gently pressing the two ends together. The nearest end is then to be closed with a cork, and the thickened portion, having been strongly heated, is to be distended into a bulb by blowing into the tube at the other end, during its rotation in the mouth. In making a pipette, a strong capillary termination should be first formed, and the bulb afterwards blown. The suction orifice may be everted or not at pleasure. The successful blowing of even small bulbs will not be found easy save after considerable practice.

(16.) Connections of tubing and apparatus are made in various ways. Two pieces of tubing of the same diameter may be sealed together, but the operation requires some amount of skill for its performance. The two ends, well adapted and by preference slightly everted, should be heated to softening in the Bunsen or blowpipe flame, and then brought steadily into contact, taking care that the edges exactly coincide. The junction has then to be heated for some time, constantly rotating the tube, and alternately pushing and pulling the free ends, one of which should be stopped with a cork, and the other occasionally blown into, so as to maintain a proper calibre. A piece of wide may be joined to a piece of smaller tubing, by first drawing out the former to a point, and then cutting it across just where its diameter coincides with that of the narrow tube. The two can then be joined together as if originally of equal size. In this way funnel-tubes may be made.

Glass tubes of more or less similar diameter are best connected by means of a short piece of vulcanite tubing, the internal diameter of which should be rather less than the exterior diameter of the glass tubes, so as to grasp them firmly by its contraction without requiring to be tied. In the absence of vulcanite, small connectors may be made of sheet india-rubber, a piece of which of the required size is to be gently warmed, and wrapped round a glass rod or tube; when its opposite edges, having been cut obliquely so as to overlap, and firmly pressed together with the thumb-nails, will, if freshly cut and perfectly clean, adhere thoroughly. The removal of the caoutchouc connector, from the tube or rod

on which it has been made, may be facilitated by first moistening the rod in the mouth or afterwards dipping it in water. Glass tubes may often be advantageously connected by a considerable length of vulcanite tubing, which can be closed at will anywhere in its course by pressing it together with a clamp of some kind, or even by tying it tightly with string. A stiff union of two tubes may be made by wrapping a piece of well-soaked bladder or parchment paper several times round their opposed ends, and allowing it to dry on. Or they may be first connected by a piece of vulcanite tubing, and then stiffened by tying on one or two wooden splints made out of lucifer matches. In this way an ordinary funnel may be readily converted into a funnel-tube. An extempore funnel-tube may also be formed by merely resting a small funnel in the suction-orifice of a plain straight pipette (fig. 9 a).

A small tube may be adapted to a considerably larger one, or to the neck of a flask or bottle, by means of a well-fitting perforated cork, the size of which can be readily reduced to any required extent by rasping and filing. A sound cork of slightly conical shape, and of such a size as to require some little force for its insertion, having been softened by pressure between the fingers or rolling under the foot, should be pierced by thrusting the point of a rat-tail file half way through it from each end, and then right through. The hole so made must be enlarged by filing until of a size to fit the tube tightly. When two or more holes have to be made in the same cork, care must be taken to have them all parallel, and as equidistant as possible from one another and from the outside of the cork. For making the necessary perforations variously-sized brass tubes with cutting edges, known as cork-borers, may be used with advantage instead of files. The holes made with them should always be rather too small at first, so as to require some little enlargement by filing. By means of cork-borers, very admirable substitutes for corks may be cut out of solid vulcanite. The end of a tube to be inserted through a perforation should always be rounded off by carefully heating it to redness, so that it may not cut or tear the bore, through which

it is, after cooling, to be gradually thrust with a screw-like motion. If fitting very tightly, it may advantageously be greased before its insertion. The cork, with its tube or tubes, is next to be fitted to the neck of the flask or bottle by direct pressure with the fingers and by gentle screwing. That the junction is air-tight may be ascertained by sucking out or blowing into the closed apparatus, and noticing with the tongue whether any exhaustion, or by the ear whether any compression, is produced. Lutes should be avoided as much as possible, but sealing-wax, varnish, white lead, and linseed meal paste are sometimes useful to stop a leak. Occasionally two pieces of tubing of more or less similar diameter are connected by being each of them inserted half-way through the opposite ends of a long perforated cork.

(17.) By means of glass tubing, vulcanite connectors, and perforated corks, together with flasks, bottles, or test-tubes, the student may construct for himself a variety of useful apparatus,

Fig. 10.



such, for instance, as the drop-bottle (fig. 10). From this, when surrounded by the warm hand and inverted, the water or other contained liquid issues in a succession of drops; but when forcibly blown into and quickly inverted, a jet of water springs forth with considerable but gradually decreasing force. This bottle, which should never be

more than half filled, is convenient for moistening substances, and for washing down into filters or evaporating dishes the frequently adhering contents of inverted tubes and beakers.

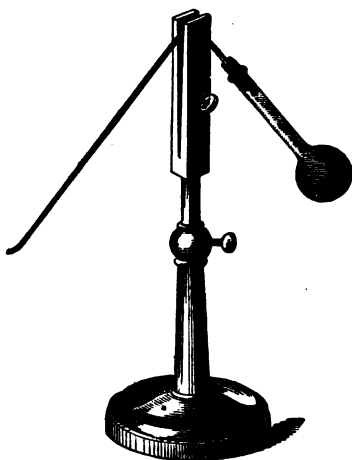
The wash-bottle also (fig. 11), when held in an upright position and blown into through the short tube, furnishes a fine strong jet of water suitable for washing precipitates; while, upon simple inversion, it delivers a coarse stream of water through the blow-tube, serving to dilute solutions, fill test-tubes, &c. The blow-tube may conveniently be made with a vulcanite joint or mouthpiece, so as to allow some freedom of movement during its use. The jet also may be attached by a cork joint, whereby it can be pointed in any direction. It is convenient to have several wash-bottles, one for cold distilled water, one made out of a flat-bottomed flask for boiling water, one of smaller size for alcohol, &c. The neck of the hot-water flask should be bound round with list or something of the kind, so as to allow of its being handled. The blow-tube of the alcohol flask may be temporarily closed by a short length of vulcanite, one end of which has been stopped with a bit of glass rod.

Retorts suitable for the generation of gases are readily made by adapting a bent tube and cork to a small flask or test-tube, as in fig. 12. But when the gas is liberable without the application

Fig. 11.

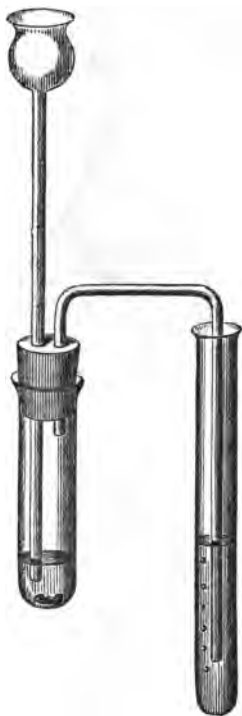


Fig. 12.



of heat, a small phial, of such a shape as to stand firmly on the table, may be used instead of a flask. The addition of a funnel-tube is useful when fresh liquid has to be supplied from time to time in order to maintain the effervescence. The bottom of this

Fig. 13.



tube must dip under the liquid in the retort or bottle, so as to be cut off from the liberated gas.

An arrangement of the kind shown in fig. 13 is useful for testing the nature of a gas generated by the action of a known liquid, usually an acid, upon some unknown substance. The acid is poured upon the other substance through the tube-funnel, and the gas conducted by the bent tube into the test solution, where its effects, if any, are observable.

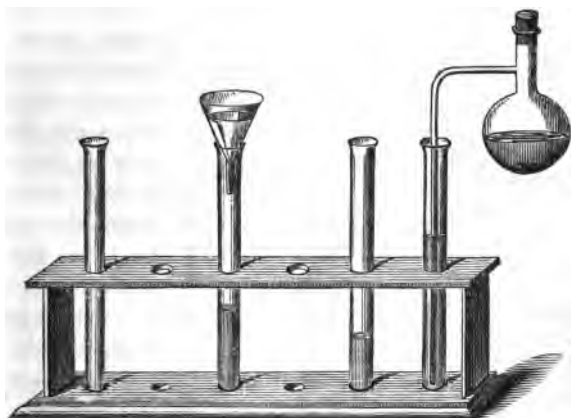
Gases often require to be purified by acting on them with certain liquid or solid reagents capable of retaining their different impurities, of which aqueous vapour is by far the most common. The solid reagent, divided into small pieces, is usually contained in a glass tube, either straight or bent in the form of the letter U,—a little cotton wool or tow being interposed between it and the perforated corks of the tube. The liquid reagent is either absorbed into pieces of pumice

or other porous solid contained in a U-tube, or the current of gas is allowed to bubble up through the liquid contained in a two-necked bottle, or in a wide-mouthed bottle, or in a U-tube, according to circumstances. Most insoluble gases, when required in a pure state, are first washed by their transmission through

water, and then dried, if necessary, by being passed over chloride of calcium or pumice soaked in oil of vitriol.

An apparatus for evolving sulphuretted hydrogen gas is indispensable to the analyst. For ordinary testing the generating bulb shown in fig. 14 is very convenient. Four or five small lumps of sulphide of iron, not larger than peas, are slid down the neck into the bulb, and covered to some depth with water. Sulphuric acid is then added little by little, until a brisk effe-

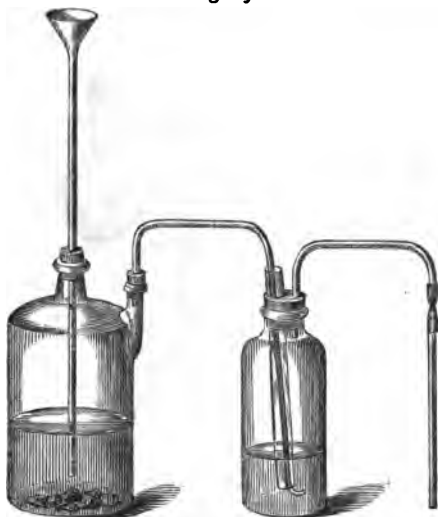
Fig. 14.



vescence is produced, which usually happens when the acid measures about one-tenth part of the water previously introduced. The mouth is then closed by a tightly fitting cork, or even by the thumb, and the evolved gas transmitted through the solution to be examined. In the absence of the above-described bulb, the arrangement shown on page 26 may be employed. The gas from either apparatus may be washed, if necessary, by transmission through a little water contained in a second generating bulb, or in a test-tube, as shown in fig. 42. But when a continuous supply of washed sulphuretted hydrogen is required, as in some toxicological experiments, a different arrangement is preferable. The gas is developed in a Wolfe's bottle, into one neck

of which there passes a tube-funnel, and from the other a delivery-tube bent twice at right angles, which dips through a wider tube into a second bottle charged with a very dilute solution of potash, and furnished with a delivery-tube conveying the gas into the solution to be precipitated, as in fig. 15. But contrivances of

Fig. 15.



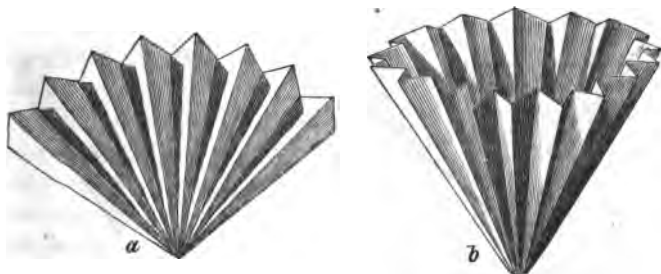
this kind may be infinitely varied, according to the fancy of the operator.

(18.) Filtration is performed in order to separate a mechanically mixed liquid and solid, with a view to obtain the clear liquid which passes through the filter, or the suspended solid which is retained on the filter, or both liquid and solid apart from one another. For analytical purposes, a fine, thin, white

blotting-paper is employed as the ordinary filtering medium. Boiling water should not dissolve anything from it, and when burnt it should leave scarcely any appreciable ash. It is sold either in sheets, or preferably cut into circles of various sizes. A round or square piece of the paper is folded, in one or other of two ways, into the form of a cone, which, unless very small, should rest in a funnel of glass or fine porcelain. When the object of filtration is merely to clarify the liquid, and especially when such liquid is at all viscid, or requires to be very quickly filtered, as often happens with a hot saturated solution, a ribbed filter is employed. The mode of folding this filter is not easily described, though

very easily learnt from demonstration. A circle or square of paper is folded first into halves, then into quarters, and each of the two double quarters again into quarters, all the creases being made on *the same side* of the doubled paper; each sector is next to be divided into two by a crease down the middle made on *the opposite side* of the still doubled paper. At this stage the filter assumes the form of a child's fan (fig. 16 *a*), and in the

Fig. 16.



event of a square of paper having been used, the projecting ends may be cut off while the fan is closed. The doubled halves are now for the first time separated, which may be facilitated by blowing on to the edge of the paper, when a deeply ribbed cone will be produced, consisting everywhere of alternate internal and external angles, except at two opposite places where two external angles will be found together, between each of which a subsequent fold must be made, so as to produce an internal angle between them. The filter is now completed, and when gently opened out has the form shown in fig. 16 *b*. The different creases should be made very sharply at the circumference, but indistinctly at the centre of the paper, for fear of weakening it too much.

But when the chief object of filtration is to collect the suspended matter, most usually a precipitate especially thrown down, a plain filter is much to be preferred. A piece of paper is folded into halves and then into quarters, when it will have the outline of an isosceles triangle, with two straight and one curved side if folded from a circle, or with three straight sides if folded

diagonally from a square, in which case the base must be cut round, as shown in fig. 17 *a*. The filter is then opened out, leaving three thicknesses on one side and one thickness on the other, so as to form a smooth cone (fig. 17 *b*), which is carefully fitted into a funnel in such a manner as to be well supported all round. The funnel should be rather larger than the filter, so as to project somewhat beyond it, while the filter itself should always

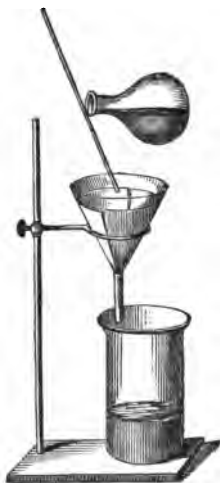
Fig. 17.



project beyond the contained liquid. Where it is necessary to employ a double filter, the two should be folded separately, and so arranged in the funnel that the three thicknesses of the one may correspond to the one thickness of the other. The outside filter is often made very small, so as merely to support the bottom of the other. A filter should always be wetted before receiving the mixture to be filtered. This is necessary in order to swell the paper and thereby close its pores, which are otherwise apt to become choked with the precipitate, if indeed some of it be not carried through by the rapid imbibition which at first takes place. When the liquid passes through the paper at all turbid, it should be returned once or twice into the filter, after which it will generally come through clear. It is usually advisable to let the mixture to be filtered subside a little before beginning its filtration. The comparatively clear liquid is then poured off into the filter, and, when it has run through, the thick sediment added separately. The interior of the tube, test-glass or beaker, which contained the mixture, is next to be washed down by a small forcible stream from the wash-bottle, and the rinsings

poured on to the deposit left in the filter by the draining away of the previously added liquid. Lastly, by means of a wash-bottle, any deposit extending up the sides of the filter is washed down to its centre, so that the entire precipitate may be collected into as small a bulk as possible, for further treatment. It is often necessary, moreover, to wash a precipitate thoroughly, so as to free it from every trace of soluble matter. This is done by projecting water upon it several times by means of the wash-bottle, and allowing the bulk of each addition to filter away before repeating the process, until, on evaporating down a few drops of the filtered washings, no residue whatever is left upon the slip of glass or platinum foil. The stream of water must not be too forcible, for fear of making a hole in the filter or causing a spurting of its contents. Moreover, in pouring into an empty filter, the liquid should be directed along the side and not immediately upon the point of the filter, which is its most unprotected and consequently weakest part; while the filtered liquid if received in a wide beaker or evaporating dish, should be made to run along the side of the glass or dish, so as to avoid spurting. Again, in pouring from one vessel into another,—from a beaker or wide-mouthed flask into a funnel for instance,—a glass rod should be applied to the lip of the flask or beaker, as shown in fig. 18, not only to direct the course of the liquid into the funnel, but also to prevent any of it being spilt by running over the side of the delivering vessel. This running of liquid over the side may also be avoided by greasing that part of the lip which is poured from, with a little tallow or spermaceti ointment.

Fig. 18.



When it is necessary to remove a wet precipitate from the filter on which it has been collected, the well-drained filter, carefully removed from its funnel, may be spread out on two or three folds of bibulous paper to absorb superfluous moisture, and the precipitate be then carefully scraped off from it by a spatula of ivory, platinum, or steel; or the drained filter may be spread out on one side of the funnel, and its contents washed quickly down by a forcible stream from the wash bottle; or, by means of a glass rod, a hole may be made in the bottom of a filter *in situ*, and its contents washed through; or the precipitate may be dissolved off the filter by causing some solvent, usually an acid, to pass through it several times, preferably at a boiling temperature.

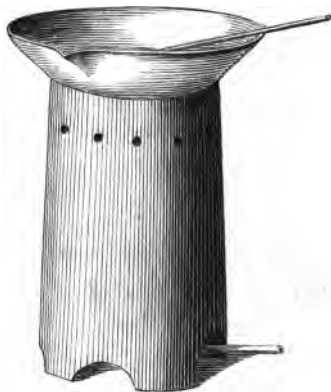
Decantation often furnishes a ready substitute for filtration in cases where the solid part of a mixture has subsided to the bottom of the precipitating glass. The speedy and complete subsidence of a freshly formed precipitate may often be effected by violently shaking up the mixture for a few minutes in a closed vessel, when, after standing at rest for a little while, the clear supernatant liquor may be poured off with a steady hand, or be withdrawn by a syphon or pipette. In using a pipette great care must be taken not to allow any of the liquid once sucked up to descend again upon the sediment so as to disturb it. A deposit from which the supernatant liquid has been removed by some form of decantation may be washed by pouring water on to it, stirring it well up, and letting it again subside for a second decantation, and so on. A thin layer of liquid overlying a deposit may often be sucked up very completely by a coil of bibulous paper, introduced with care so as not to disturb the deposit.

Syphons and pipettes are useful not only for separating a supernatant liquid from a deposited solid, but also for separating two strata of different liquids from one another.

(19.) Apparatus to be heated over the several burners already described, may, according to its nature, be supported in various ways. It may rest on the ring, or be held by the clamp, of a retort-stand; or an independent upright clamp or tripod stand may be employed. The tripod is often replaced with advantage by a

jacket of clay, or metal, surrounding the burner, and so preventing draughts, as shown in fig. 19. Iron triangles also to rest on the top of the jacket, tripod, or retort-ring, are in constant requisition. A triangle of iron wire, sheathed with three pieces of tobacco-pipe, is useful for supporting small porcelain crucibles that have to be made red-hot, but the mass of the tobacco-pipe interferes with the attainment of a very high temperature. Small triangles of platinum wire are far more convenient.

Fig. 19.



Flasks and retorts may be heated at some little distance over an argand flame without the interposition of any medium. But with the gauze burner, and more particularly with the Bunsen burner, it is advisable to protect the bottom of the vessel by a piece of stout wire gauze. Beakers should be placed either on a flat iron plate or sand-bath, and the heat be raised cautiously. Porcelain evaporating dishes may be heated almost anyhow, according to circumstances; but when the flame touches the bottom of the dish some little care is necessary, both at the commencement of the operation, and when the liquid is evaporated nearly to dryness. Watch-glasses require very careful heating. They are best held by the thumb and finger over a small flame, but may be supported by forceps, wire-triangle, or special watch-glass holder. When containing liquid, to be heated by means of a sand-bath, they should not be depressed into, but just rest on, the top of the hot sand. Test tubes may be readily heated in the flame of a spirit lamp or gauze-burner. They should be held between the thumb and fingers, and be constantly shaken from side to side, especially during boiling, to prevent any

sudden expulsion of the contained liquid, with which they should not be more than half filled. When the boiling has to be long continued, a piece of paper or cloth may be bound or twisted

Fig. 20.

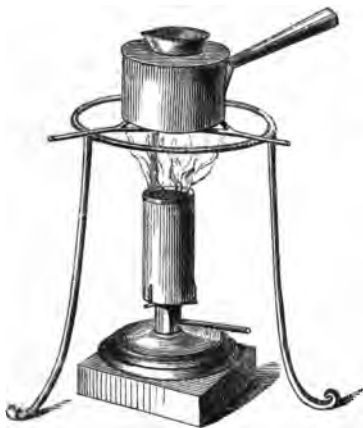


round the upper part of the tube, so as to protect the fingers. Test tube holders are rarely of much use.

Small capsules and crucibles, of platinum or Berlin ware, may be heated to redness over an argand (fig. 20), or to full redness over a Bunsen burner, or by the blowpipe flame. They may be held either with forceps or on triangles of wire or tobacco-pipe. When a strong heat is required, they should be surrounded with a small jacket of metal or clay.

Sand-baths are usually formed of sheet iron. Some are made rather deep to receive flasks and retorts, others nearly flat for

Fig. 21.



beakers. The sand, which must not be heaped above the level of the liquid to be heated, should be of uniform size and rather coarse. An iron plate, or trellis of thick iron wire, forms an excellent substitute for a flat sand-bath.

The water-bath is used for heating substances to a temperature not exceeding 100° C. A small saucepan, with an evaporating dish for a cover, forms a capital makeshift. The saucepan lid may be replaced by a series of broad rings of tin plate, having

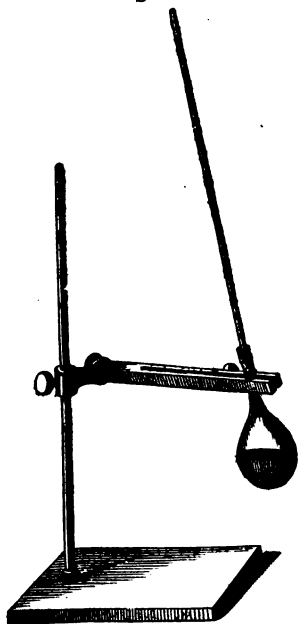
apertures of different sizes to support small evaporating dishes (fig. 21), capsules, watch-glasses, necks of flasks, &c. Beakers, flasks, and retorts, to be heated in a water-bath, should not be allowed to touch the bottom of the bath, but should rest on a piece of tow or folded cloth, as well to avoid danger from bumping as to prevent the communication of an increased temperature by contact with the heated metal.

(20.) Heat is applied to liquids in order to warm, evaporate, boil, or distil them. Evaporations are performed on slips of flat glass, or on watch-glasses, or evaporating dishes. The crystalline forms of various salts may often be recognised by dissolving a grain or so of the salt in a drop or two of water on a glass slip, evaporating until a solid margin appears, and setting aside to crystallise. The residue may then be examined by a lens, or under the low power of a microscope. The evaporation of a small quantity of liquid, contained in a watch-glass for instance, may often be promoted by gently blowing on its surface for a little while. A dish in which evaporation is taking place (fig. 19) may be loosely covered with a piece of filtering paper, stiffened by a glass strip passed through and across it; and care should be taken not to allow the liquid to boil. In evaporating to dryness, it is well to have the heat lowered as the process approaches completion, and the residue, if considerable, kept constantly stirred.

Ebullition is conducted on a small scale in test tubes, and on a larger scale in flasks,—Florence oil flasks being among the best and cheapest that can be employed. A flask of cold liquid, held over a naked flame, quickly becomes covered with a deposit of moisture, which it is advisable to wipe off once or twice. Ebullition sometimes takes place intermittently, and with considerable jerking or bumping. This may often be prevented by introducing a coil of platinum wire or piece of tobacco pipe into the liquid, either before boiling or after cooling down a degree or two. If introduced during boiling, it is apt to produce a violent rush of vapour. It is occasionally useful to adapt a long upright tube to

the mouth of a flask (fig. 22) in which a liquid has been heated, so that any vapour given off may be condensed in the tube, and flow back again into the flask.

Fig. 22.



Distillation is usually conducted in an apparatus composed of a boiler, condenser, and receiver. On a small scale the boiler is represented by a glass retort, or flask with its bent tube, the receiver by a test tube, flask or bottle, and the condenser by a long glass tube placed between the retort and receiving flask, and surrounded either by a constantly changing layer of water, as in Liebig's condenser, or by a piece of blotting-paper kept moist by the constant dripping of water. Very often a separate condensing tube is dispensed with, and the long neck either of the retort or receiving flask alone employed. Fig.

23 shows a very simple arrangement of this description. The liquid is boiled in a long-necked retort, and the distillate conducted by means of a bent adapter into an upright bottle. A twist of thoroughly wetted tow, or lamp-cotton, is tied somewhat tightly round the retort-neck, at about an inch above the point where it enters the perforated cork of the adapter, and the ends of the twist allowed to hang down for two or three inches. A piece of filtering paper of suitable size and shape to embrace almost the entire circumference of the neck, and reach from just above the twist of tow almost to the curve of the retort, is moistened with water, laid upon the retort-neck and bent round it so as to adhere closely. A second piece of filtering paper is at first folded by means of

three creases into four strips and then placed over the other, so that the flaps being bent down, its middle portion will form a

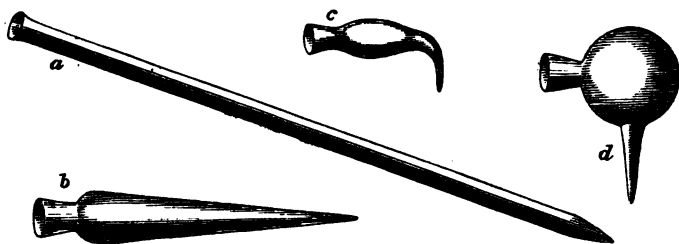
Fig. 23.



small channel along the top of the retort-neck. It should be about two-thirds the length of the other, and not reach so high up. By this arrangement the water constantly dropping out of a partly plugged funnel from the height of half an inch or so on to the surface of the inner paper, at a little distance from its upper extremity, is conveyed along the channel, spread uniformly over the retort-neck and drained away by the twist of tow. If the short length of neck intervening between the twist and adapter be dry at the beginning, it will continue so throughout the experiment. For further security, however, it may be marked with a ring of grease. Adapters are of all shapes and sizes. Such a one as that shown in the woodcut is easily made out of a piece of tubing or broken retort neck. Others, intended to act more or less as condensers, are represented in fig. 24. Another convenient form of distilling apparatus is shown in fig. 40. The boiler consists of a round flask, from which a long wide bent tube dips through a perforated cork into a Florence flask resting in a basin of water.

The cork must either not fit tightly or have a slit cut in it, so as to permit the escape of any uncondensed vapour. The neck and upper surface of the receiving flask should be surrounded with

Fig. 24.



filtering paper, on to which water should drop from a funnel. A bend of sheet lead may be placed over this flask so as to sink it in the water.

One of the simplest varieties of Liebig's condenser is shown in

Fig. 25.

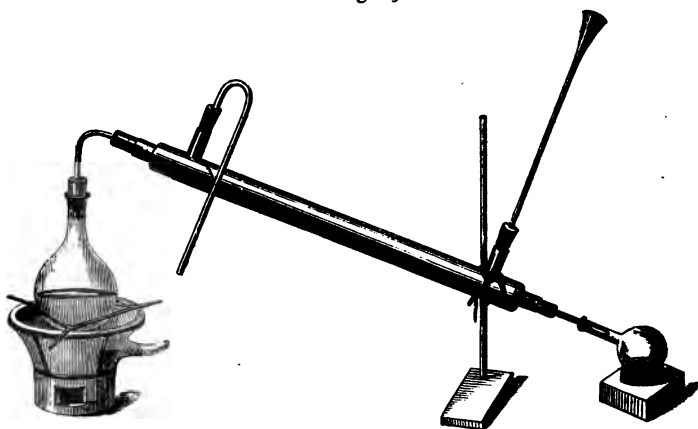


figure 25. It is merely a cylinder of tin-plate, having four slightly conical tubular apertures—two in the same plane with each other at either side of the cylindrical surface, and two oppo-

site each other in the terminal circular planes. By means of well-fitting perforated corks a funnel tube of tin-plate or glass is inserted into the distant horizontal aperture, and an exit tube of tin or glass into the other, while the distilling tube extends through the axis of the cylinder. The hot contents of the distilling tube pass downwards, becoming colder and colder in their descent, while the stream of cold water entering the condenser through the funnel passes upwards, becoming hotter and hotter in its ascent until it escapes at the overflow pipe. The condenser may be fastened to a retort-stand with string or wire, as in the figure, or may be supported by a clamp of some kind. Liebig's condensers are made in every variety of form, size, material, and construction,—some of them being provided with special supports, which allow them to be heightened or lowered at will, and placed at any desired inclination.

(21.) Heat is applied to solids in order to warm, dry, ignite, fuse, or volatilise them. Any tube, flask, or retort, the interior of which cannot be reached by the fingers, should, after thorough cleaning,* be rinsed once or twice with distilled water, and then drained as dry as possible by means of draining pegs or some other mode of support. It should next be warmed carefully over a gas flame or in front of a fire, and the hot moist air sucked out of it from time to time by the aid of a long tube reaching into it for some

Fig. 26.

considerable distance, as shown in fig.

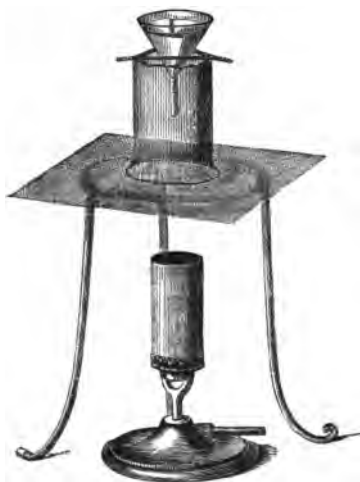
26. Narrow glass tubing is



* There is seldom much difficulty in cleaning laboratory glass from any stain or dirtiness, when it is not of long standing, by means of cold or hot water and dilute or strong acids or alkalis, aided by extempore brushes made of moist tow dipped in sand and fastened on to thick pieces of wire, or by tube-brushes made specially for the purpose. Flasks, retorts, &c., may often be very efficiently cleaned by shaking them up somewhat violently, after the introduction of a little water and a few pieces of soft paper or rag. The interior of narrow glass tubing is best cleaned by pushing a piece of moist filtering paper through it.

dried by heating some length of it over a gas burner or spirit lamp, and simultaneously sucking air through it with the mouth. In the absence of an air- or water-oven, reduction-tubes and similar small pieces of apparatus may be dried by heating them on a sand-bath standing over a burner; or preferably on a flat iron plate, which in many other cases also may be advantageously substituted for a sand-bath. Moist powders of various kinds may be dried on a water-bath, or sand-bath, or by ignition over an argand burner, &c., according to circumstances. A washed precipitate retained in its filter and funnel may often be quickly

Fig. 27.



dried by supporting the funnel on a broken beaker or short lamp-glass standing upon a trellis of iron wire, underneath which a small gas-flame is kept burning (fig. 27). Or the filter may be supported over a heated iron plate by means of a small tripod-stand, easily made out of copper-wire. When nearly dry, the filter with its contents may be removed from the funnel and placed in a water-bath; or the washed precipitate and filter may be removed carefully from the funnel, pressed

gently between folds of blotting-paper, and placed at once in the water-bath. At moderate temperatures, drying over oil of vitriol in the exhausted receiver of an air-pump takes place with considerable rapidity. Moreover, a shallow air-pump-jar standing on a plate of ground glass over a dish of oil of vitriol or quicklime, forms a very convenient chamber in which all sorts of bodies may be dried and kept dry.

Animal solids are frequently subjected to ignition in order to

burn off their organic, and leave behind their mineral matter, or ash. The tissue, &c., may be first carbonised in small portions at a time in a thin Berlin capsule or crucible, heated over a gauze burner, in some place where the empyreumatic vapour, &c., can be readily got rid of. The resulting charcoal should then be pulverised, and the powder heated steadily for some hours in a shallow platinum capsule, or on a tray of platinum foil, supported over an argand flame, when the charcoal will gradually burn away, and a white or greyish ash be left. The temperature should never exceed that of dull redness, as otherwise the ash, save that of blood, is apt to fuse over the remaining charcoal, and so prevent its combustion. The operation is much facilitated by protecting the capsule from draughts, and particularly by placing over, but not immediately upon it, a cover of thin platinum foil. Carbonate of sodium and other fluxes employed in testing, are often heated to dull redness over an argand flame just before being used. Moreover, in quantitative analysis, filters and their contained precipitates have constantly to be burnt, with a view to getting rid of the filter-paper and leaving the precipitate in a state fit for being weighed. The ignitions made on charcoal or platinum wire in the course of blowpipe testing will be presently described.

Independently of the many fusions made in the course of blowpipe testing, others on a somewhat larger, though still very small scale, have occasionally to be performed by the student. There are, for instance, a few substances which require to be fused with carbonate of sodium or potassium, either alone or mixed with some other reagent, before they can be brought into a state of solution, and so identified by ordinary analytical processes. The insoluble substance is usually incorporated with three or four times its bulk of a mixture of carbonate of sodium with carbonate of potassium, or in some cases with either the nitrate or cyanide of potassium, and heated to thorough fusion over a Bunsen or blowpipe flame, in a platinum or porcelain capsule, or in a small iron spoon. In making these fusions, it is most important that both the substance and flux be well dried, very finely powdered, and intimately commixed. The capsule or crucible should be heated

at first very gradually, but ultimately to the highest attainable temperature.

The only volatilisations which the student will be called upon to perform are made in narrow glass tubes,—open at both ends when a current of air is required to act upon the heated substance, or open at one end only when a simple sublimation is intended. Powdered substances of various kinds may be introduced into narrow tubes open at both ends, by first placing a suitable quantity of the powder in a gutter of stiff glazed paper, pushing this gutter with its contents into the tube held horizontally, then inverting the tube and gutter, and, lastly, withdrawing the gutter while still inverted. The same method may sometimes be used with closed subliming tubes, so as to avoid soiling their interiors, but is unnecessary when both substance and tube are thoroughly dry. The tube, whether open or closed at one end, should be made of hard glass, and be heated in the flame of a spirit lamp or Bunsen burner.

(22.) Although quantitative analysis does not come within the scope of this work, yet a few words on weighing and measuring may not form an inappropriate addition to the foregoing remarks on chemical manipulation. The general adoption of the French metrical system, of which the gramme is the unit of weight, and the cubic centimetre or bulk of a gramme of water at its greatest density the most usual unit of measure, is highly desirable; but in default of this, the English decimal system, of which the standard grain is the unit of weight, and the bulk of a grain of water, at 62° F., the unit of measure, may be employed. In the chemical laboratory we dispense altogether with the use of ounces, drams, &c., and speak only of so many grammes and cubic centimetres, or grains and grain-measures. Mr. Griffin takes the bulk of seven grains of water as his unit of measure, which he terms a septem, so that 1000 septems are equal to one decigallon, or to the bulk of a pound of water. The use of this decimal division of the gallon is often very convenient, and quite compatible with that of the grain-measure, the septem and grain-measure standing to one another in the simple relation of 7 to 1,

as shown in the following table. The figures with a dot over them are inexact.

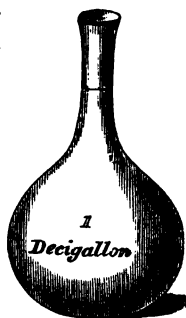
	Decigallons	Septems	Grain-measures
Gallon	10,000	70,000
Decigallon or pound	1,000	7,000
Quart	2·50	2,500	17,500
Pint	1·25	1,250	8,750
Fluid ounce	·0625	62·5	437·5
Cubic foot	62·321	62,321	436,247·4
Cubic inch	36·065	252·458
Litre	2·204	2,204·6	15,432·6
Cubic centimetre	2·2	15·4

One litre, or kilogramme of water-measure, equals 1·76 pints, or 61·027 cubic inches. One cubic centimetre, or gramme of water-measure, equals ·061 cubic inch.

One decigallon or pound of water-measure equals ·4535 litre, or 453·5 cubic centimetres, or 0·0160 cubic foot, or 27·727 cubic inches, or 16 fluid ounces.

Fig. 28.

Some measures are made to contain or deliver a definite quantity of liquid. Others are graduated so that any indefinite quantity delivered may be afterwards read off. A flat-bottomed and somewhat narrow-necked flask, having a horizontal scratch across its neck marking the height to which it should be filled, forms a very convenient measure of the former kind (fig. 28). It is easy to select a couple of such flasks, which shall measure either a decigallon and half-decigallon respectively, or a litre and half-litre, &c., &c. A pipette of the form shown in fig. 29 is also a very useful instrument of this class. It is filled by carefully sucking up liquid to a level somewhat above the mark on its stem, and quickly closing its upper orifice by the finger. Then by



relaxing the pressure of the finger, the contained liquid is allowed to issue drop by drop until its height corresponds exactly with the

scratch, when the finger is again pressed down; after which, on removing the

finger, the measured quantity of liquid is allowed to flow out. A pipette of this kind is so constructed that when filled up to the mark on its stem it delivers exactly the indicated quantity of liquid, irrespective of what adheres to the interior of its elongated bulb. The last drop should be gently blown out while the point of the pipette is in contact with the inside of the receiving vessel. It is well to have a couple of such pipettes delivering either 100 septems and 10 septems, or 50 and 5 cubic centimetres respectively. For measuring indefinite quantities a tall narrow cylinder (fig. 30), graduated into divisions of 10 septems, or 5 c. c. each, is most convenient. The ordinary ounce measures of the apothecary are very unsatisfactory instruments. Their graduation is seldom accurate and always difficult to read off exactly, on account of the expanded conical form of the glass. Where smaller quantities have to be delivered and their volumes afterwards noticed, a Bink's burette or a graduated straight pipette may be employed. The burette (fig. 31 a) should be held near its upper end, with its mouth guarded by the thumb or forefinger, and its beak pointed not directly but obliquely downwards, so that the side of the beak may be inclined to the horizontal plane at a somewhat acute angle. The orifice of the beak should be well greased with tallow



Fig. 29.

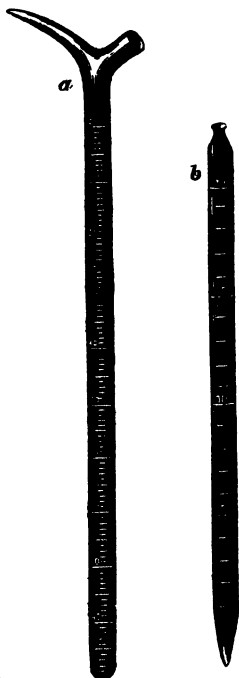


Fig. 30.

or spermaceti ointment, and any liquid remaining in it be sucked

down, both at the beginning and end of the experiment. Moreover, the burette must always be allowed to stand at rest for a minute or two before observing the height of the contained liquid. The graduation should be from above to below, as in the woodcut.

Fig. 31.



The graduated pipette (fig. 31 *b*) is filled by suction, the contained liquid adjusted to a proper height, the quantity required allowed to flow out, and the level of the remainder finally read off. Inasmuch as the conical extremity of this kind of pipette rarely delivers its contents exactly, it is better not to have it included in the graduation. Whether the graduation of the pipette is from above to below, or from below to above, does not much matter; for some purposes the one, and for some purposes the other mode being most convenient.

In reading off the height of a liquid its upper surface should be brought as nearly as possible to a level with the eye. This surface in most liquids contained in glass vessels, will be found more or less deeply concave according to the diametric smallness of the column, &c.; but, in all cases, the bottom of the curve is taken as the true level, and, in measuring definite quantities, must be made to coincide exactly with the mark. By right all measurements should be taken at mean temperature, namely, 15.5° C. (60° F.), but the expansions and contractions of aqueous liquids within the ordinary ranges of temperature are so slight that, in most cases, they may be safely disregarded.

A pair of the best description of dispensing scales suspended

from a fixed support makes a very useful balance for ordinary work. A set of accurate grain weights ranging from 0.05, or 0.1 grain to 1000 grains should be provided, and also a supplementary short pan for taking specific gravities. The beam should turn freely by an addition of 0.1 grain, even when the pans are each loaded with a weight of twelve or fifteen hundred grains. It is well to employ habitually the left-hand pan for the substance, and the right-hand pan for the weights, which should be always handled by pincers, and not by the fingers. It is sometimes necessary to weigh out definite quantities of a substance such as 20, 50, or 100 grains; but it is better in most cases to take an indefinite quantity and then ascertain its weight, exactly as in weighing any particular specimen, the resultant of an experiment for instance. When taking the weight of an indefinite quantity, much time will be saved by trying the weights not at random but in a definite order, always taking in succession the weight next above or below in the series that particular weight which was last found too little or too much.

It is convenient to provide counterpoises of sheet lead for balancing exactly the several watch-glasses, capsules, crucibles, bottles, &c., used for retaining a substance while being weighed; or their respective weights may be ascertained and scratched upon them. When a definite quantity of substance has to be transferred from the watch-glass or capsule on which it has been weighed into some other vessel, the frequently adhering residue may be washed off by a jet of water, or be gently brushed off by a camel's-hair pencil. But where an indefinite quantity has been taken, it is better to reweigh the watch-glass or capsule and to subtract the weight of adhering residue from the original weight of substance. Or, what comes to the same thing, some portion of a weighed quantity may be transferred, and its amount ascertained by noticing the loss sustained by the originally weighed quantity.

Substances to be accurately weighed must always be first brought to an uniform condition of dryness, inasmuch as a greater or less degree of dryness may cause considerable variation in the

weight of a body at different times. The substance may be dried in vacuo over oil of vitriol, or in a water-bath, and its weight determined from time to time until it becomes constant; or, in some cases, the substance may be heated at once to dull redness and then weighed,—not, however, until thoroughly cooled, as by resting the containing crucible on a piece of metal, for instance, otherwise an ascending current of heated air will be set up, which will diminish its apparent weight appreciably. Hygro-metric substances must be weighed in covered crucibles, or between a pair of ground watch-glasses held together by a clip, or in stoppered bottles made expressly for the purpose, or in short wide test tubes. They may be cooled under a glass jar standing over oil of vitriol.

(23.) By specific gravity is understood the weight of a unit of volume, or, what comes to the same thing, the comparative weights of equal bulks of different bodies. In this country water at the temperature of 15.5° C. (60° F.) is taken as the standard to which the specific gravities of liquids and solids are usually referred, and its sp. gr. considered either as 1.0 or 1000 according to circumstances.

In order to determine the specific gravity of a liquid, a small flask or bottle of known weight is selected, which, when accurately filled to a certain point, contains a known weight of pure water at 15.5° . This flask is filled with the liquid whose sp. gr. is required, and weighed, when, after deducting the known weight of the flask, the residuary weight of contained liquid is compared with the known weight of the same bulk of water, according to the proportion :

Wt. of water W' : wt. of liquid W :: sp. gr. of water : sp. gr. of liquid ;

that is, according to the equation,

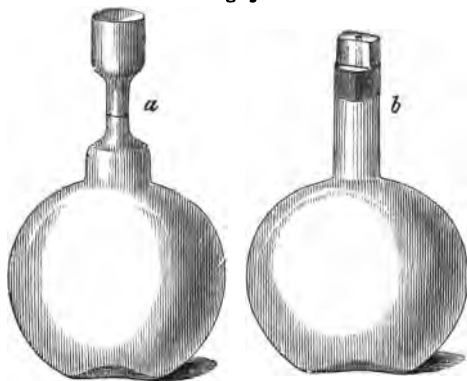
$$\text{Sp. gr. of liquid} = \frac{W \times 1000}{W'}.$$

Thus, supposing the flask to be what is called a five-hundred

grain flask, but to contain in reality 499.7 grains of water, and say 459.8 grains of proof spirit, then the sp. gr. of the spirit will be $\frac{459.8}{499.7} = 920$.

Specific gravity flasks or bottles are made of various shapes, one of the most convenient being that employed by Regnault and shown in fig. 32 *a*, for which, however, any flat-bottomed narrow-

Fig. 32.

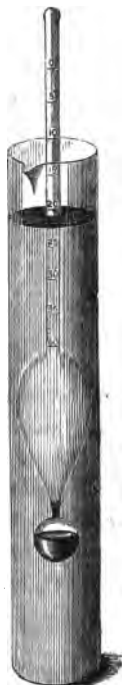


necked flask may be substituted. It is to be filled up to the mark on its neck with the liquid, previously brought to the temperature of 15.5° by immersion in cooled or warmed water, and weighed. For taking the specific gravities of very volatile liquids

this kind of flask is provided with a solid stopper. The most usual form of sp. gr. flask is shown in fig. 32 *b*; when used, it is nearly filled with the liquid to be examined, a tube thermometer introduced, and the whole placed in a vessel of cooled or warmed water until the contained liquid has acquired a temperature of 15.5° . The thermometer is then removed, and the flask filled to the brim with more of the liquid previously brought to mean temperature, the perforated stopper inserted, whereby the excess of liquid is thrust out, and the exterior of the flask thoroughly dried with a cloth, care being taken to avoid any communication of heat from the hand or elsewhere. As a rule sp. gr. flasks should be made perfectly dry before being filled, and by preference be rinsed out with some of the liquid under examination.

The hydrometer (fig. 33) is useful for taking the specific gravities of different liquids, where rapidity rather than accuracy of determination is required. It is merely an upright float, weighted below, and having a narrow graduated stem above, so as to sink to a greater or less extent in liquids of different densities. Inasmuch as the bulk of any liquid displaced by a floating body is equal in weight to the floating body, it is obvious that equal weights of different liquids will differ in volume according to the depth to which the hydrometer sinks in them; or, in other words, the heights to which different liquids rise on the stem will be inversely as their specific gravities. Some hydrometers, those used in the examination of urine, for instance, are so graduated that their degrees express the specific gravities directly. But in commerce various artificial scales are preferred, that of Twaddell in particular being very generally used in this country. The degree of gravity marked on Twaddell's scale has to be multiplied by 5, and the product added to 1000, to give the actual specific gravity.

Fig. 33.



The most usual method of taking the specific gravity of a solid body heavier than water, consists in weighing it first in air, or theoretically in a vacuum, and afterwards in pure water at 15.5, when the ratio of the difference in the weighings to the weight in air will give the sp. gr., according to the proportion :

Dif. in wgs. $W - W'$: wt. in air W :: sp. gr. of water : sp. gr. of solid ;
that is, according to the equation

$$\text{Sp. gr. of solid} = \frac{W \times 1.000}{W - W'}.$$

For by a well-known principle in hydrostatics, the apparent loss

of weight which a body experiences when immersed in a liquid is identical with the weight of an equal bulk of that liquid.*

In order to weigh a solid body in water it must be attached by a horse-hair to the hook of the sp. gr. pan, as shown in fig. 34,



and have its surface thoroughly wetted with a soft brush, so that, when immersed in the water, there shall be no adhering air bubbles. When the solid is soluble in water it must be immersed in alcohol, petroleum, or some other liquid of which the specific gravity has been previously ascertained, and the calculation made as before, substituting the sp. gr. of the liquid employed for the sp. gr. of water. When the solid is lighter than water it must, after its weight in air has been taken, be attached to some heavy body sufficient to sink it, a piece of lead, for instance, and the weight of the solid

and piece of lead in water and of the piece of lead alone in water ascertained. The weight of the volume of water displaced by the light solid will equal the weight of the light solid in air W , plus the buoyancy it imparts to the immersed lead, as measured by the difference between the weight of the lead in water w , and the conjoint weight of the lead and light body in water w' . Hence we have the proportion :

Wt. of water $W + w - w'$: wt. of solid W :: sp. gr. of water : sp. gr. of solid ;
or the equation :

$$\text{Sp. gr. of light solid} = \frac{W \times 1.000}{W + w - w'}.$$

* Relying on this principle, the specific gravities of different liquids may be ascertained by immersing some solid of known weight in each of them, and comparing the losses in weight which it experiences by the upward pressures of the different liquids, equal in each instance to the weight of the bulk of liquid displaced by it.

When the substance to be examined is in the pulverulent state, its specific gravity may be ascertained by means of a sp. gr. flask. A quantity of the powder is introduced into the flask and weighed. It is then thoroughly wetted with water or some other liquid, of which a further quantity is afterwards added, so as to fill up the flask to the mark on its neck; when the weighing is repeated. The difference between the weight of liquid which the flask ordinarily holds, and the weight of liquid added to fill up the flask after the introduction of the dry powder, gives the weight of the bulk of liquid displaced by the powder, the ratio of which to the weight of the powder gives the specific gravity.

CHAPTER II.

ANALYTICAL CHEMISTRY.

(24.) THE object of this part of the course is to make the student practically acquainted with the chemical properties of such bodies as are of the most importance, and of the most common occurrence.

Of all chemical compounds, those known as salts will most frequently present themselves to his notice.

Sulphate of iron and chloride of sodium may be taken as the types of simple salts. The sulphuric acid and chlorine are termed the electro-negative or acid constituents, or, more shortly, the acids; the iron and sodium the electro-positive or basic constituents, or, more shortly, the bases.

In testing the substances distributed for examination, each of which should contain but one basic and one acid constituent, the student may first of all confine himself to the bases; subsequently he must examine both for bases and acids. He will have to pursue the following course of operations:—

I. To examine the dry substance before the blowpipe. (Pars. 25 and 26.)

II. To make a solution of the substance in water or acid. (Pars. 27 and 28.)

III. To ascertain to which group the base of his substance belongs. (Pars. 30, 35, and 41.)

IV. To identify the particular member of the group with which he is dealing. (Tables I. II. and III.)

V. To realise the special reactions described under the head of his particular base.

VI. To identify the acid constituent of his substance. (Tables IV. and V.)

VII. To realise the special reactions described under the head of his particular acid.

As regards V. and VII., the student must remember that the simple discovery of the base and acid of his salt is of much less importance than the thorough verification of all their described properties.

§ I.—BLOWPIPE EXAMINATION.

(25.) A small shallow hole having been made on a piece of charcoal, the student should put into it a little of the substance under examination, a piece about the size of a mustard-seed, for instance, or as much powder as will rest on the point of a penknife. He must then heat the substance on its charcoal support before the blowpipe, and notice what effects, if any, are produced. It is often advisable to moisten a pulverulent substance with water before submitting it to the blowpipe flame, so as to make it cohere and remain on the charcoal. The substance may also be heated with advantage in a subliming tube open at both ends; whereby corroborative, and sometimes even primary, evidence of its composition is obtainable, especially when it happens to be wholly or partly volatile. In this way ammoniacal and sulphurous acid gases may be recognised by their respective smells; deposits of acid or alkaline water, by their reaction to test-paper; and sublimes of sulphur, arsenic, mercury, and ammonium-salts, by their appearance and behaviour.

The following are the most important general effects observable upon heating a substance on charcoal before the blowpipe.

All hydrated salts give off their water of crystallisation, some with intumescence, as borax; others with decrepitation, as gypsum. Many anhydrous salts also, as chloride of sodium,

for instance, decrepitate from the expulsion of water retained mechanically within their crystals. Most hydrated salts, when first heated, fuse in their water of crystallisation, and then solidify, whether or not susceptible of again fusing at a higher temperature, (vide par. 26).

Most compounds of the heavy metals become darker when heated, either permanently through decomposition, or temporarily through some altered relation to light. Zinc compounds acquire a deep greenish-yellow by the action of heat, and peroxide of tin a pale brownish-yellow.

Certain metallic compounds, more especially those of silver, lead, and bismuth, are quickly brought to the metallic state when heated on charcoal in the reducing blowpipe flame, (vide par. 26).

Some substances, whether or not inflammable, leave a black carbonaceous residue, capable of being burnt away by prolonged ignition. This charring indicates the presence of organic matter, —an organic acid or base, for instance, either free or in combination.

Many bodies evolve a more or less marked smell when heated. Thus sulphur and many sulphides give off sulphurous anhydride. Tartaric and benzoic acids, and their respective salts, evolve characteristic empyreumatic odours. The fixed organic bases, and some salts of ammonia and organic bases, give off ammoniacal vapour; while arsenic compounds, when in contact with ignited charcoal, give rise to a peculiar garlic-like smell.

Sometimes the heated substance volatilises wholly or in great measure, usually with evolution of visible fumes, (vide par. 26).

In a few cases the red-hot charcoal undergoes a rapid combustion where it comes in contact with the heated compound. This deflagration indicates the probable presence of a nitrate or chlorate.

(26.) As regards their more special behaviour before the blowpipe, metallic compounds may be classified into those which are volatile (α), those which leave a white permanently fusible

residue (β), those which leave a white infusible residue (γ), those which are reducible to the metallic state (δ), and those which give a colouration to the borax bead (ϵ), as shown in the following scheme:—

BLOWPIPE EXAMINATION

a. Volatile	β . White and Fusible	γ . White and Infusible	δ . Reducible	ϵ . Colour borax bead
AMMONIUM	SODIUM	ZINC	SILVER	CHROMIUM
MERCURY	POTASSIUM	ALUMINUM	TIN	<i>Green</i>
Salts	Salts	MAGNESIUM	LEAD	MANGANESE
ARSENIC	CALCIUM	CALCIUM	BISMUTH	<i>Amethyst</i>
Oxides	STRONTIUM	STRONTIUM	ANTIMONY	IRON
Sulphides	BARIUM	BARIUM	CADMIUM	<i>Yellow-brown</i>
ANTIMONY	Chlorides	Salts	<i>Give incrustations.</i>	COBALT
Teroxide	<i>Colour blow-pipe flame.</i>	SILICIO	ZINC γ	<i>Deep-blue</i>
OXALIC ACID		STANNIC	MERCURY α	NICKEL
		ANTIMONIC	ARSENIC α	<i>Reddish</i>
		Oxides	COPPER ϵ	COPPER
				<i>Pale-blue</i>
				<i>In oxidising flame</i>

a. The ordinary compounds of ammonium and mercury are readily volatile. Phosphate of ammonium, however, leaves a fused residue of phosphoric acid, which requires a strong heat for its dissipation. The oxides and sulphides of arsenic also are readily volatile; the teroxide of antimony somewhat less so. Oxalic acid melts and effervesces during its disappearance without furnishing much incrustation. The compounds of mercury, arsenic, and antimony, and many ammonium-salts, deposit incrustations or sublimates upon the cold part of the charcoal. Similar sublimates are procurable by heating the substance in a sealed or open subliming tube.

The various bodies which volatilise when heated on charcoal

may often be satisfactorily identified by a few rough tests, such as the following :—

Ammonium salts, when warmed with potash, evolve ammonia, recognisable by its odour and alkaline reaction. Mercury salts, save the black and red sulphide, are turned black by sulphide of ammonium. Mixed with a large excess of carbonate of sodium, and heated in a reduction tube, they afford a sublimate of mercurial globules. The oxides (white) and sulphides (orange or yellow) of arsenic dissolve in sulphide of ammonium to form a yellowish liquid, which on evaporation to dryness leaves a bright yellow residue. Mixed with soda-flux, and heated in a reduction tube, they furnish sublimed crusts of metallic arsenic. Teroxide of antimony dissolves in sulphide of ammonium, and the liquid, when evaporated down, leaves a deep orange residue. It is, moreover, easily reducible before the blowpipe (δ). Oxalic acid, when moistened with water, manifests a strongly acid reaction, and effervesces with peroxide of manganese.

β . The ordinary salts of sodium and potassium fuse at a red heat, many of them into almost watery liquids, which are absorbed by the porous charcoal. During their ignition, the salts of sodium impart a bright yellow, and those of potassium a violet colouration to the blowpipe flame. These colours are best seen by taking up a minute quantity of the salt upon the end of a platinum wire, and heating it at the point of the blowpipe flame. When a potassium salt is contaminated with even a small proportion of any sodium salt, the violet colour of its flame is liable to be concealed by the strong yellow colour which the sodium salt produces. But the proper potassium colour may be readily seen by looking at the flame through a piece of smalt glass, which cuts off the yellow sodium rays.

Unlike most salts of the alkaline earth-metals, the chlorides of calcium, strontium, and barium are fusible in the blowpipe flame. That of calcium fuses very readily, but the other two chlorides are much less readily fusible than are the majority of sodium and potassium salts. When strongly ignited on fine

platinum wire, chloride of barium imparts to the blowpipe flame a marked apple green, chloride of strontium a deep crimson, and chloride of calcium an orange-red colour. The platinum wire used for testing by the blowpipe flame should be moderately thin, to allow of its being strongly heated. It may be held directly in the fingers, platinum being a bad conductor of heat. Should the end to be used impart any colour to the flame, from contamination either with soda, derived from the perspiration of the fingers, or with the residues of former experiments, it must be alternately dipped in hydrochloric acid, and strongly heated in the blowpipe flame until all colouration ceases to appear.

The chlorides of the alkaline earth-metals, and most salts of the alkali metals, fuse into colourless transparent liquids, whereby they are distinguished from the fusible compounds of the heavy metals, which yield coloured or opaque beads. Moreover, the temporary melting of hydrated salts in their water of crystallisation must not be confounded with the permanent fusion at a red heat which characterises the above-mentioned classes of salts.

γ. Compounds of zinc, aluminum, and magnesium, with nearly all calcium, strontium, and barium salts, whether or not undergoing a preliminary aqueous fusion, leave after strong ignition a white infusible residue; while the stannic, silicic, and antimonie oxides are from the first infusible. During strong ignition the aluminum residue manifests an intense white incandescence, the zinc residue a deep greenish-yellow, and the stannic and antimonie oxides a pale brownish-yellow colour. In any case the white or yellowish-white infusible residue may be moistened with solution of nitrate of cobalt, and again strongly heated, whereby characteristic colourations are produced with compounds of zinc, aluminum, and magnesium, and less definite colourations with the remainder. The zinc residue acquires a fine green, the aluminum residue a bright blue, and the magnesium residue a very faint pink colour. When the residue is pulverulent, it may be moistened with sulphuric acid and re-ignited before being heated with

nitrate of cobalt, more especially in order to bring out the magnesian colour. It must be borne in mind that nitrate of cobalt also imparts a blue colour to many fused phosphates, borates, and silicates, which, however cannot be confounded with the blue given to an infusible aluminum residue.

The other infusible residues receive a less distinct colour by ignition with nitrate of cobalt, those of calcium and strontium becoming grey, that of barium reddish-grey, those of the stannic and silicic anhydrides bluish-grey, and that of antimony greenish-grey; but these substances may be further distinguished by other means,—calcium, strontium, and barium by their tinting the blowpipe flame,—antimony and tin by their reducibility, the former with, and the latter without, any incrustation (δ),—and silica by its behaviour with fused carbonate of sodium. In order to test an infusible compound for calcium, strontium, or barium, a small quantity, taken up on the end of a fine platinum wire, should be moistened with hydrochloric acid, and ignited for some time in the hottest part of the blowpipe flame, when the characteristic colour due to the volatilisation of each metal will be clearly brought out, save indeed with their respective sulphates. Silica may be identified by making a small loop at the end of a platinum wire, and fusing upon it a minute quantity of carbonate of sodium, which will thus form a white bead, transparent when hot, opaque when cold: in this fused bead, silica, when strongly heated, will dissolve with effervescence, and, if in sufficient quantity, render it permanently transparent.

δ . Compounds of silver, tin, lead, bismuth, antimony, and cadmium, often assume the metallic state, or produce characteristic incrustations, when merely heated on charcoal in the reducing blowpipe flame; but their behaviour when heated with a flux of carbonate of sodium, mixed or not with cyanide of potassium, is more satisfactory. Carbonate of sodium alone will answer the purpose, but the addition thereto of one-fourth of its weight of cyanide of potassium often assists the reduction very greatly, and is not in any case disadvantageous, save when the

specimen itself deflagrates upon charcoal from the presence of a nitrate or chlorate, in which case a slight explosion results from the reaction of the cyanide and oxisalt.

The specimen having been intimately mixed with five or six times its bulk of flux, a small portion of the resulting powder, sometimes moistened with water so as to make it cohere, is heated strongly on charcoal. The mixed mass should fuse readily before the blowpipe, so that any minute globules of reduced metal may run together. If not readily fusible, a fresh mixture must be taken with a larger proportion of flux. When the reduced metal volatilises at the temperature employed, its vapour becomes oxidised outside the flame, and is deposited upon the charcoal as a more or less abundant, white or coloured, incrustation. Silver gives no incrustation, and tin scarcely any: lead gives a yellow, and bismuth a brownish-yellow, incrustation; while antimony gives an abundant easily volatile bluish-white, and cadmium a comparatively fixed brown-red incrustation. Metallic antimony vaporises rapidly; while cadmium is so volatile that its reduction and vaporisation are simultaneous, wherefore no globule, but only an incrustation, is producible with it. Zinc compounds, heated with reducing flux, behave in this respect like cadmium, furnishing no globule of metal, but only an incrustation, which is yellow when hot, white when cold; but zinc will have been previously detected by its reaction with nitrate of cobalt (γ). Reduced arsenic and mercury are so volatile that they can only be obtained in the form of sublimates by performing the reduction in tubes as already described (α).

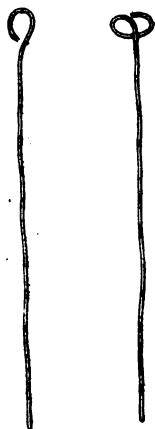
The different metals usually identified by their reduction on charcoal, exhibit the following characters:—Silver yields a bead of white, moderately hard and malleable metal, with no incrustation. Tin, which is less easily reducible, yields a bead of white malleable metal, softer than silver, with very slight, if any, incrustation. Lead yields a bead of soft bluish-white metal, with yellow incrustation; bismuth, a bead of brittle yellowish-white metal, with brownish-yellow incrustation; antimony, a bead of brittle bluish-white metal, with an abundant

bluish-white incrustation; and cadmium, no metallic bead, but only a reddish-brown incrustation.

In addition, copper, iron, nickel, and cobalt are reducible, though they do not afford incrustations, and are not easily obtainable in the form of beads. But on crushing the fused mass in a mortar, and washing away the lighter portions, copper may often be recognised in the form of red spangles, and iron, nickel, and cobalt as heavy powders affected by the magnet.

c. Compounds of those metals which tinge the borax bead usually leave dark-coloured infusible residues when heated alone on charcoal. In testing by the borax bead, a piece of platinum wire is bent into a single or double loop, as shown of the actual

Fig. 35.



size in fig. 35, the loop dipped into powdered borax, and the adhering borax heated to redness, when it first undergoes a highly characteristic intumescence, and afterwards, when more strongly heated, sinks into a colourless transparent bead. To this bead is then attached a minute quantity of the substance under examination, and the whole strongly heated in the blowpipe flame, when in some cases the fused bead dissolves the specimen, and thereby acquires a more or less definite colour, the depth of which may be increased by adding more of the specimen and again heating strongly.

The metals iron and copper, which form two classes of salts, also form beads of two colours. Thus in the oxidising flame we have a blue cupric and a yellow-brown ferric bead, while in the reducing flame we have a cuprous bead of an almost colourless or opaque reddish aspect, and a ferrous bead of a sea-green colour. The chromium bead has an emerald green, and the cobalt bead a sapphire blue colour. Manganese, when free from iron, imparts an amethystine tint, and nickel a deep sherry hue, which becomes amethystine when the bead is heated with a

fragment of nitre. The borax may be replaced by microcosmic salt, or even by ordinary glass. In these several reagents we have, after ignition, an excess of melted boric, phosphoric, or silicic anhydride, which at the temperature of the blowpipe flame combines with the various metallic oxides to form coloured fusible salts.

§ II.—SOLUTION AND PRECIPITATION.

(27.) Having made his examination in the dry way, by means of the blowpipe, the student must next bring his substance, by some means or other, into a state of solution, so that he may submit it to the action of liquid reagents. As a general rule, the substance to be dissolved should be in a finely divided state. This is particularly necessary in the case of bodies which are with difficulty soluble, such as many native oxides, sulphides, &c. Any substance having a decided colour, a hard structure, and an opaque aspect, whether earthy or lustrous, ought always to be pulverised very finely before being treated with solvents. The solution of the body under examination should be effected by preference in water; but, if insoluble in water, it may be acted upon with hydrochloric acid, or with nitro-muriatic acid, or with nitric acid.

A small portion of the powdered substance is to be placed in a test-tube, a moderate quantity of water* added, the whole agitated, and heated over a spirit or gas-flame. While heating, the tube should receive an occasional jerk, to facilitate mixture and avoid the sudden escape of vapour. If the substance, by this treatment, is obviously dissolved, the clear solution, filtered if necessary from any insoluble portions, can be submitted at once to the action of reagents. If the substance, however, is not obviously dissolved, a few drops of the liquid may be filtered on to a glass slip and gently evaporated to dryness. Should any

* By water is always meant pure or distilled water; but clean rain water may sometimes be employed as a substitute.

definite amount of residue remain upon the glass, the whole mixture must be thrown upon a filter, and the tests applied to the clear filtrate. There are many substances which, unless taken in very small quantity, do not disappear perceptibly when boiled in water, but yet are sufficiently soluble to afford an aqueous solution that can be successfully tested. Should a mere trace of residue, or none at all, be left upon the glass slip, as much of the water as possible is to be poured away from the insoluble substance, and replaced little by little with hydrochloric acid, warming between each addition. Should any obvious action occur, more hydrochloric acid may be added, if necessary, and the whole heated for some time until an available solution is formed. Should there be no obvious action, nitric acid must be added in the proportion of about one-fourth of the hydrochloric acid previously used, and heat again applied. By one or other of these means a solution will generally be effected. There are, indeed, a few substances which dissolve in nitric, but neither in hydrochloric nor in nitro-hydrochloric acid. There are also some substances which are quite insoluble in any ordinary menstruum; the consideration of these, however, is deferred for the present. The solution of the substance, whether in water or acid, to which no reagent has been added, is called in the tables and elsewhere *the original solution*. The freshly-made acid solution should generally be diluted somewhat freely with water, and filtered if necessary. The acidity, neutrality, or alkalinity of the aqueous solution should be ascertained by means of test-paper.

(28.) Different reagents are next to be added to the original solution in the order and manner described in paragraphs 30, 35, and 41, and in the Tables I. II. III. and V. These reagents produce in the solution certain effects, which are characteristic of the various substances dissolved. The effect most usually produced by a liquid reagent is to cause a *precipitate* or solid deposit of some insoluble compound of the substance sought for. Hence the formation or non-formation of a particular precipitate usually proves the presence or absence of a particular base or acid in the solution under examination. Precipitates differ much in their

colour, consistency, rapidity of formation, and solubility in different liquids, whence the student must make himself familiar with their various aspects and habitudes. As regards aspect, he must notice whether a precipitate is dense, crystalline, clotty, gelatinous, opaque, transparent, coloured or colourless, &c. As regards habitude, he will find that crystalline precipitates, unless thrown down from concentrated solutions, do not usually appear at once, but only after some little time. Their immediate formation, however, may be often determined by rubbing the liquid against the inside of the containing vessel with a glass rod. Again, many precipitates are characterised by their solubility in an excess of the precipitant, or in some other reagent.

Reagents and solvents should always be added gradually, except when special direction is given to the contrary. This rule is of great importance, and applies equally to the formation and solution of precipitates; in the latter case, the mixture should be agitated between each addition of the solvent. Many characteristic effects are occasionally overlooked through a neglect of this rule. The student must also bear in mind, when directed to employ an excess of any particular reagent or solvent, that every minute quantity more than sufficient to produce the desired effect is an excess.

In the tables, the word *dissolved* placed at the head of a column signifies either that the substances written under it have not been precipitated at all, or that, having been precipitated, they are now redissolved by an excess of the reagent, in any case that they remain in solution.

EXAMINATION FOR BASIC GROUPS.

(29.) The bases are divided into three principal groups, as follows :—

I.	II.	III.
TIN	NICKEL	BARIUM
ARSENIC	COBALT	STRONTIUM
ANTIMONY	MANGANESE	CALCIUM
BISMUTH	IRON	MAGNESIUM
MERCURY	CHROMIUM	POTASSIUM
LEAD	ALUMINUM	SODIUM
SILVER	ZINC	AMMONIUM
COPPER		
CADMIUM		

The first object of the student must be to ascertain the group to which the base of the salt under his examination belongs.

The members of the first group are precipitated from their acid solutions by *sulphydric acid*. The members of the second group are not precipitated from their acid solutions by *sulphydric acid*, but are precipitated from their neutral solutions by *sulphide of ammonium*. The members of the third group are precipitated neither by *sulphydric acid* nor by *sulphide of ammonium*.

Having ascertained to which particular group the base of his salt belongs, he will proceed according to the directions of the table pertaining to that group, in order to identify the individual member thereof with which he is dealing. Should, therefore, *sulphydric acid* produce a precipitate, in an acidified solution, the student will proceed according to Table I. Should it produce no obvious precipitate, he will render the solution nearly neutral by *ammonia*, and then add *sulphide of ammonium*. Should this reagent produce a precipitate he will proceed according to Table II. Should no precipitate be produced by either of the above reagents, he will proceed according to Table III.

§ III.—EXAMINATION FOR BASES OF GROUP I.

(30.) To recognise the presence of some member of this group by means of *sulphuretted hydrogen* or *sulphydric acid*, the solution to be tested should be moderately acid. In solutions which are too acid, sulphuretted hydrogen may not give any precipitate, despite the presence of a member of the group; and in solutions which are neutral or alkaline, it may give a precipitate even in the absence of every member of the group; inasmuch as sulphuretted hydrogen precipitates some members of the second group from their neutral or alkaline solutions.

AQUEOUS SOLUTIONS. These must consequently be acidulated before being treated with sulphuretted hydrogen. A few drops of either nitric or hydrochloric acid will answer the purpose, but the use of the latter acid is generally preferable. The addition of hydrochloric acid, however, sometimes produces a permanent white precipitate, in which case the presence of silver, or lead, or mercury is indicated. Solutions of silver invariably yield a precipitate with hydrochloric acid, solutions of lead and mercury only under certain conditions. But in the event of hydrochloric acid producing a precipitate, it will suffice for the student to distinguish between the above three metals, without following out the directions of the general table for the group (page 68.)

The **SILVER** precipitate is soluble in excess of ammonia.

The **MERCURY** precipitate is turned black by excess of ammonia.

The **LEAD** precipitate is unaffected by ammonia, but is soluble in boiling water; and, on cooling, is deposited therefrom in crystalline needles.

These three precipitates are blackened by sulphuretted hydrogen, and are not produced by nitric acid, properties distinguishing them from all other precipitates which hydrochloric acid occasionally produces.

The acidification of a solution of tartar-emetic, with either hydrochloric or nitric acid, is attended with the production of a white turbidity, which, however, disappears on gently warming

the liquid with a little more acid. Moreover, the acidification, by either hydrochloric or nitric acid, of various alkaline solutions, not unfrequently gives rise to whitish precipitates, which sometimes disappear in an excess of acid, and, at other times, remain. Among those which are permanent, the principal are sulphur, from the decomposition of an alkaline persulphide, &c.; silica, from the decomposition of an alkaline silicate; and boric acid from the decomposition of an alkaline borate; but this last precipitate is readily soluble in boiling water.

SOLUTIONS IN ACID. When the solution of the original substance has been made in an acid, it is important to get rid of any great excess of acid; or, at any rate, to reduce its activity.

a. By mere dilution with water. It is generally advisable to dilute somewhat considerably solutions which have been made by means of an acid.

β. By evaporation. The solution may be evaporated down to a small bulk, and then be diluted with water. This process is especially necessary when the solution has been made with nitro-muriatic acid.

γ. By neutralisation with ammonia. When a large quantity of acid has been employed to effect the solution of a substance, it is occasionally useful to neutralise some of the excess of acid with ammonia.

The acid solution of the substance, whether or not evaporated down, or partly neutralised, should, after dilution with water, be perfectly bright. If not bright, it must be rendered so by filtration.

The addition of water to an acid solution sometimes produces an obvious white precipitate, in which case the dilution should be very slight or be dispensed with altogether. The formation of a white precipitate on the addition of water indicates the presence of **ANTIMONY** or **BISMUTH**. The precipitate produced in solutions of the former metal is dissolved by tartaric acid and turned of an orange colour by sulphuretted hydrogen; while that produced in solutions of the latter metal is not dissolved by tartaric acid, and is turned black by sulphuretted hydrogen. Water does not

invariably cause a precipitate in solutions containing antimony or bismuth, but in the event of a precipitate being produced, it will suffice to distinguish between the above two metals without proceeding according to the general table for the group (page 68).

The acidified solution of the substance in water, or the diluted solution of the substance in acid, is to be treated with sulphuretted hydrogen. It may be sufficient to add sulphuretted hydrogen water to the solution, but it is always preferable to use a current of the gas itself. The production of a coloured precipitate is indicative of the presence of some member of the first group; in which case, the gas should be passed into the liquid until it smells permanently even after agitation. A little water should next be added, and the whole well shaken or stirred, to promote the subsequent subsidence of the precipitate, which, on setting the tube aside for a few minutes, will soon collect at the bottom. The supernatant liquid may then be poured off, and the precipitate treated according to the directions of Table I. β .

Sulphuretted hydrogen, when added to certain solutions, not containing any member of the first group, sometimes produces a more or less considerable yellowish-white turbidity, due to a liberation of finely divided sulphur, effected by some per-oxidated or per-chlorinetted compound, thus:



In yellow solutions, this white turbidity often appears decidedly yellow, from the colour of the liquid through which it is seen. When, simultaneously with the liberation of sulphur, a brownish-yellow solution becomes paler or colourless, the presence of a per-salt of iron may be generally inferred; but when it becomes of a marked green colour the presence of chromic acid is indicated.

TABLE I.

(31.) Examination of a solution containing some one member of the first group; namely, TIN, ARSENIC, ANTIMONY, BISMUTH, SILVER, MERCURY, LEAD, COPPER, or CADMIUM; all of which metals are precipitated from their acid solutions by *Sulphuretted hydrogen gas* (a), or its solution in water.

β. Having treated the not too acid solution of the substance with excess of sulphuretted hydrogen, and poured off or filtered off the supernatant liquid, warm the precipitate with some solution of *Sulphide of Ammonium*.

If the precipitated sulphide be of

TIN (*protosalt*), brown,

TIN (*persalt*), yellow,

ARSENIC, yellow,

ANTIMONY, orange,

it will *dissolve* entirely, and on addition of Hydrochloric Acid to the resulting solution will be reprecipitated.

TIN, always yellow

ARSENIC, yellow

ANTIMONY, orange

γ. Boil the precipitate thrown down by sulphuretted hydrogen with strong *Hydrochloric Acid*.

Undissolved

ARSENIC

Dissolved

TIN

ANTIMONY.

If the precipitated sulphide be of

BISMUTH

SILVER

MERCURY

LEAD

COPPER

CADMIUM, yellow,

it will remain *undissolved*.

δ. Add *Potash* to a portion of the original solution: in any case a precipitate will be produced characterised as follows:—

LEAD, white, *soluble* in excess of the reagent, unaffected by ammonia.

MERCURY (*protosalt*), black, unaffected by ammonia.

MERCURY (*persalt*), yellow, turned white by ammonia.

BISMUTH, white, unaffected by ammonia.

COPPER, blue

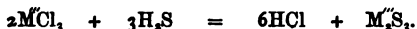
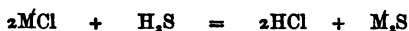
CADMIUM, white

SILVER, brown

} soluble in ammonia.

Insoluble in excess of potash.

(32.) *a. Sulphydric acid, or sulphuretted hydrogen, reacts with the salts belonging to this group to form metallic sulphides, as shown by the following typical equations, in which M stands for an atom of metal :*



In this manner the salts of perissad metals, as silver Ag' , arsenic As''' , antimony Sb''' , and bismuth Bi''' , are decomposed by sulphuretted hydrogen.



In this manner the salts of artiad metals, as lead Pb'' , mercury Hg'' , copper Cu'' , cadmium Cd'' , tin (stannosum) Sn'' , and tin (stannicum) Sn''' are decomposed by sulphuretted hydrogen.

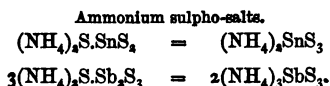
The sulphides thus produced differ much from one another as regards their solubility in mineral acids. They are all completely dissolved by nitro-muriatic acid; except that of silver, which is converted into insoluble chloride of silver; and that of lead, which is converted partly into the sparingly soluble chloride, partly into the insoluble sulphate of lead, owing to an oxidation of its constituent sulphur.

Hot nitric acid dissolves the sulphides of arsenic, bismuth, silver, copper, and cadmium, but has no appreciable action on the sulphides of mercury. It converts the sulphides of tin and antimony into their insoluble oxides or anhydrides, SnO_2 and Sb_2O_3 , respectively. When slightly diluted, it dissolves sulphide of lead completely, but otherwise it converts a portion of it into insoluble sulphate of lead. The action of nitric acid upon the sulphides is generally attended with a separation of sulphur, which, on boiling, gradually assumes the form of melted globules.

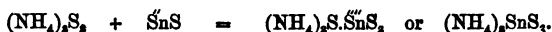
Strong hydrochloric acid at a boiling temperature has no action on the sulphides of arsenic and mercury. It converts the sulphides of silver and lead into their insoluble or sparingly

soluble chlorides, and dissolves the remaining sulphides of the group with greater or less facility.

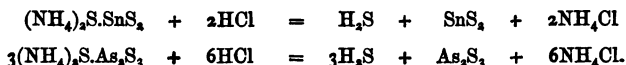
(33.) β . The disulphide of tin, and the trisulphides of arsenic and antimony, unite with the sulphides of alkali-metal to form soluble sulphur-salts corresponding to the well-known oxygen salts, thus:



Hence these sulphides are distinguished from the remainder by their solubility in *sulphide of ammonium*.* Protosulphide of tin is not soluble in pure colourless sulphide or sulphhydrate of ammonium; but it is soluble in the ordinary yellow solution of the persulphide, whereby it becomes converted into the above-described compound of disulphide of tin, thus:



On the addition of hydrochloric acid to any of these sulpho-salts, they are decomposed with reprecipitation of their respective sulphides, thus:



It is advisable not to dissolve the sulphides of this sub-section in an unnecessarily large quantity of yellow sulphide of ammonium, lest the subsequent addition of hydrochloric acid should separate so great a quantity of whitish sulphur as to conceal the colour of the reprecipitated sulphides.

γ . Sulphide of arsenic is distinguishable from the sulphides of tin and antimony by its insolubility even in boiling hydrochloric acid; and by its solubility in a warm solution of sesquicarbonate of ammonia.

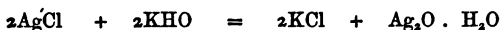
* Sulphide of copper is quite insoluble in the sulphides of sodium and potassium; but is slightly soluble in sulphide of ammonium, especially when it contains, as it usually does, some free ammonia.

Sulphide of antimony and persulphide of tin are distinguishable from one another by their difference in colour. When pure they dissolve completely in hydrochloric acid; but as usually obtained they often contain excess of sulphur, which remains undissolved. Upon evaporating down their hydrochloric acid solutions to a small bulk, stannic and antimonious chlorides are obtained respectively. The former chloride does not have its transparency affected by dilution, neither does the diluted liquid yield any deposit upon a surface of metallic tin: and again, an acid solution of chloride of tin, in which a small fragment of zinc has been dissolved, gives with corrosive sublimate a white precipitate of calomel, gradually becoming grey from its conversion into metallic mercury (vide par. 34).

The latter chloride is generally rendered opaque by diluting its solution, which again becomes clear on the addition of tartaric acid; while the diluted liquid yields an abundant black deposit of pulverulent antimony upon a surface of metallic tin. Moreover, chloride of antimony reacts satisfactorily when examined by Marsh's or Reinsch's process.

Solid compounds of arsenic are most readily recognised by the reduction test (vide par. 66).

(34.) *δ*. Potash reacts with the salts belonging to the second section of this group, to precipitate the hydrated oxides of the respective metals, thus:



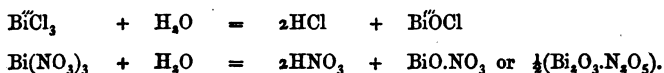
The hydrated oxides of mercury, lead, copper, and cadmium are produced according to the following general equation:



Hydrate of lead is soluble in excess of potash; the hydrates of silver, copper and cadmium are soluble in excess of ammonia; while those of bismuth and mercury are insoluble in either reagent. Independently of their behaviour with sulphuretted hydrogen and

caustic alkalis, the metals of this sub-section of the first group are characterised by the following reactions.

Bismuth solutions, unless too acid, when they must first be evaporated down, yield an opaque white precipitate on the addition of *water*, due to the formation of some insoluble basic salt, thus :



These basic salts of bismuth are insoluble in tartaric acid, and are blackened by sulphuretted hydrogen or sulphide of ammonium.

Mercury solutions yield with *protochloride of tin* a white precipitate becoming grey spontaneously, or more rapidly on the application of heat. The white precipitate is calomel, which is formed from mercurous salts by double decomposition, thus :



But it is formed from mercuric salts by reduction, thus :

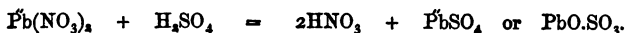


The grey deposit consists of finely divided metallic mercury, produced by an abstraction of chlorine from the calomel first precipitated :



This grey deposit, when boiled with hydrochloric acid, acquires the characteristic appearance of globules of mercury.

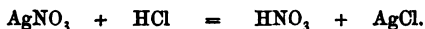
Lead solutions yield with *sulphuric acid*, or soluble sulphates, a white precipitate of sulphate of lead, insoluble in cold nitric or hydrochloric acid :



The precipitate is distinguished from the similar precipitate

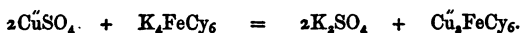
produced with barium- and strontium-salts, by its solubility in excess of potash, by its solubility in boiling hydrochloric acid, and by its becoming blackened by sulphuretted hydrogen or sulphide of ammonium.

Silver solutions yield with *hydrochloric acid* or soluble chlorides a white clotty precipitate of chloride of silver:



The precipitate is soluble in ammonia, but insoluble in the strongest nitric acid, even when boiling. It is turned of a slate-purple colour by exposure to light.

Copper solutions, even when very dilute, give with *ferrocyanide of potassium* a chocolate-red precipitate of ferrocyanide of copper, or of ferrocyanide of copper and potassium, thus:



The precipitate is turned of a pale blue colour by potash, and is then readily soluble in ammonia, forming a deep purple coloured liquid, by which properties it is distinguished from the similarly coloured ferrocyanide of uranium.

Cadmium solutions are specially recognised by the bright yellow colour and insolubility in sulphide of ammonium, of the precipitated sulphide of cadmium CdS , produced by sulphuretted hydrogen or sulphide of ammonium. Of all the sulphides of the first group of metals, sulphide of cadmium is the one most readily soluble in acids. Cadmium-salts, moreover, are readily identified by their behaviour before the blowpipe.

§ IV.—EXAMINATION FOR BASES OF GROUP II.

(35.) The members of this group are precipitated by *sulphide* or *sulphydrate of ammonium*, but are not precipitated from their acidified solutions by sulphuretted hydrogen. Inasmuch as sulphide of ammonium also precipitates most of the metals of the first group, their absence must be ascertained by the non-produc-

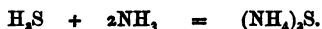
tion of a precipitate with sulphuretted hydrogen, before the reaction with sulphide of ammonium can be depended on as a general test for members of the second group.

The solution to be tested with sulphide of ammonium should be nearly neutral, but it may be slightly alkaline, or slightly acid without disadvantage. It must not, however, be decidedly acid; for if so, there may not only be no precipitate produced when some member of the group is present, but, owing to a customary impurity in the reagent, there may even be a precipitate produced when every member of the group is absent. This last occurrence is consequent upon a deposition of sulphur from the mutual decomposition of the acid solution and the reagent, quite irrespective of the presence of any metal. Pure colourless sulphide or sulphhydrate of ammonium, indeed, is not precipitated by mere acid solutions; but the yellow persulphide of ammonium, into which it becomes gradually converted, is decomposed by all acid liquids with precipitation of sulphur, thus :



The aqueous solution of a salt may be examined for members of the second group, by adding sulphide of ammonium at once; but a solution of the salt in acid must be rendered neutral, or nearly neutral, with ammonia before applying the test. The addition of even an excess of ammonia to the acid liquid is no disadvantage. It will sometimes, indeed, produce a precipitate, but the formation of a precipitate by ammonia, equally with the formation of a precipitate by sulphide of ammonium, indicates the presence of a member of the second group; though the non-production of a precipitate by ammonia does not prove the absence of all members of the group. The precipitate produced by ammonia generally differs in its character, and frequently in its appearance, from that produced by sulphide of ammonium, but the formation of a precipitate by ammonia will not interfere with the action of the more characteristic reagent for the group.

The acidulous solution of the substance which has been tested with sulphuretted hydrogen, but which has not yielded any precipitate therewith, may be examined for members of the second group by treatment with ammonia. In this case, one portion of the ammonia neutralises the excess of acid, while another portion combines with the sulphuretted hydrogen to form sulphide or sulphhydrate of ammonium, which serves to precipitate any member of the group, thus :



There are certain salts of barium, strontium, calcium, and magnesium which do not dissolve in water, but which are readily soluble in dilute mineral acids—nitric or hydrochloric, for instance—without, at the same time, undergoing any obvious decomposition. Hence, when such an acid solution is neutralised by ammonia, or by sulphide of ammonium, the salts are reprecipitated in their original condition ; so that, although the alkaline earths strictly belong to the third group, they are occasionally precipitated along with the proper members of the second. These salts are principally the fluoride of calcium—the oxalates of calcium, strontium, and barium—and the phosphates of magnesium, calcium, strontium, and barium. In Table II. they are referred to under the general term of earthy salts ; and the mode of distinguishing them from one another is described in par. 59.

TABLE II.

(36.) Examination of a solution containing some one member of the second group of bases; namely, NICKEL, COBALT, MANGANESE, IRON, CHROMIUM, ALUMINUM, or ZINC; all of which bodies are precipitated by *Sulphide of Ammonium* (a), from their neutral, or nearly neutral, solutions.*

B. Add gradually a considerable excess of aqueous *Potash* to a portion of the original solution. In any case a precipitate will be formed, which may either remain or be redissolved.

<i>Undissolved</i>		<i>Dissolved</i>	
NICKEL, pale green.		CHROME, green.	
COBALT, pale blue.		ALUMINUM, white.	
MANGANESE, white, becoming brown.		ZINC, white.	
IRON, olive-green, black, or red.			
EARTHY SALTS, white.			
<p>γ. Add to a fresh portion of the original solution, some <i>Chloride of ammonium</i> and an excess of <i>Ammonia</i>.</p>			
<i>Precipitated</i>	<i>Dissolved</i>	<i>Precipitated</i>	<i>Dissolved</i>
IRON, red	NICKEL, blue.	CHROME, green	ALUMINUM.
EARTHY SALTS, white.	COBALT, brownish pink.		ZINC.
	MANGANESE, colourless, speedily becoming brown and turbid.	To the potash solution	
		ε. Add a few drops of <i>Sulphide of Ammonium</i> .	ζ. Add excess of <i>Chloride of Ammonium</i> .
		<i>Precipitated</i> ZINC.	<i>Precipitated</i> ALUMINUM.

* To work successfully, the student must supplement the brief directions of the different tables, and more especially of this table, by the explanations and limitations of the succeeding letterpress.

(37.) *a. Sulphide of ammonium* reacts with salts pertaining to this group, to form precipitates of various characters and aspects. The precipitate is white in solutions of zinc, aluminum, and the earthy salts; black in solutions of iron, nickel, and cobalt; greenish in solutions of chrome; and buff-coloured in those of manganese. The student must not, however, attach too much importance to the colour of a precipitate, as it is a quality very liable to be interfered with by accidental circumstances. For instance, the presence of a trace of iron, occurring as an impurity, may effect a great alteration in the characteristic appearance of precipitates due to chrome, manganese, zinc, aluminum, or earthy salts, respectively, by imparting to them a black, grey, or greenish colour. From its transparency, the precipitate produced in aluminous solutions is very liable to be overlooked.

The salts of nickel, cobalt, manganese, iron and zinc are precipitated by *sulphide of ammonium*, in the form of sulphides, according to the general equation,



From their neutral solutions these metals are precipitated very imperfectly by sulphuretted hydrogen, in consequence of the formation during the reaction of hydrochloric or some other acid, in which the respective sulphides are soluble, thus:



But the sulphides of nickel, cobalt, and zinc may be precipitated completely from solutions which are acid only with acetic acid, and in which, owing to the addition of an alkaline acetate, no stronger acid than the acetic can be set free during the reaction; though even then sulphide of iron can be but partially precipitated, and sulphide of manganese not at all.

Although the sulphides of nickel and cobalt are not precipitated in the presence of hydrochloric acid, yet, when once produced, they can only be dissolved in the acid with considerable difficulty; but they are readily soluble in nitric acid. The sulphides of

zinc, iron, and manganese, however, are easily dissolved by cold hydrochloric acid, and that of manganese by acetic acid.

The sulphides of aluminum and chromium cannot be produced in the moist way. Hence the salts of these metals are precipitated by sulphide of ammonium in the form, not of sulphides, but of hydrated sesquioxides, with liberation of sulphuretted hydrogen, thus:



The earthy salts are precipitated as such by a mere neutralisation of the acid in which they were dissolved, thus (vide page 75):



(38.) β . From protosalts of nickel, cobalt, manganese, iron, and zinc, *potash* throws down the respective prothydrates, precisely as it does the prothydrates of the metals of the first group:



From the tri- or sesquisalts of iron, chromium, and aluminum, it precipitates the respective sesquihydrates, thus:—



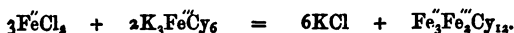
The earthy salts are precipitated as such by the neutralisation of the acid in which they were dissolved.

The appearance of the manganese precipitate is very characteristic. From being quite white it becomes rapidly brown by an absorption of atmospheric oxygen. The precipitate given by potash with perfectly pure protosalts of iron is greenish-white, but the precipitate ordinarily obtained has a dark olive-green colour, becoming ochrey-red by exposure to air: with mixed proto- and sesqui-salts, a black precipitate, and with pure sesqui-salts a red-brown precipitate is produced. The hydrates of chromium, aluminum, and zinc are completely soluble in excess

of potash, while the other precipitates are insoluble. Should the excess of potash effect an obvious solution of the precipitate at first thrown down, but the liquid at the same time not become perfectly bright, it may be filtered off from the oxide of iron, or other insoluble impurity, and tested for chromium, aluminum, and zinc (δ).

(39.) γ . *Ammonia* throws down from solutions of nickel, cobalt, manganese, iron, and earthy salts, precipitates which have respectively the same composition as those thrown down by potash. The precipitates produced in solutions of the earthy salts and in sesquisalts of iron are equally insoluble in excess of ammonia as in excess of potash; but the prothydrates of nickel, cobalt, and manganese are readily soluble, and that of iron sparingly soluble, in excess of ammonia, when chloride of ammonium or some other neutral ammoniacal salt is also present. Hence when the previous experiment with potash has shown the probable presence of a protosalt of iron, it is necessary, before adding ammonia, to boil the solution with a little nitric acid for a few minutes, so as to convert the protosalt into a sesquisalt, whereby all the iron may be precipitated as a red-brown sesquihydrate, quite insoluble in excess of ammonia.

But when the colour of the precipitate produced by potash, or some other reaction, has indicated the absence of a protosalt of iron, the boiling with nitric acid may be dispensed with. The presence of a protosalt of iron in the original solution is best shown by the addition of *ferridcyanide of potassium*, which produces therewith an abundant dark blue precipitate of ferridcyanide of iron, or Turnbull's blue:



The presence of a sesquisalt of iron is best shown by *sulphocyanate of potassium*, which produces therewith a blood-red liquid, due to the formation of sesqui-sulphocyanate of iron:



The mode of examining the earthy salts is described in par. 59.

When excess of ammonia is added to a strongly acid solution, or when chloride of ammonium and excess of ammonia are added to a nearly neutral solution of nickel, cobalt, or manganese, the precipitate at first produced is immediately redissolved.

The ammoniacal solution of nickel has a deep purple-blue colour, closely resembling that of the similarly constituted solution of copper. Potash added to the ammoniacal nickel solution throws down an apple-green precipitate of hydrate of nickel; but a very large quantity of potash may be added to the ammoniacal copper solution without disturbing its transparency, and the precipitate finally produced is for the most part blue. Moreover, ferrocyanide of potassium produces with ordinary salts of nickel a pale green, and with ordinary salts of copper a chocolate-red, precipitate of the respective ferrocyanides; and the two metals are otherwise well characterised.

The ammoniacal solution of cobalt has a brownish-pink colour, which gradually becomes darker by exposure to air, until eventually a brownish precipitate of hydrated sesquioxide of cobalt is produced. But the cobalt solution, when made with ammonia containing carbonate of ammonium, or even with carbonate of ammonium itself, has a fine pink colour, which is tolerably permanent. Cobalt compounds are best recognised by fusion with the borax bead, to which they impart a deep purple-blue colour.

The ammoniacal solution of manganese is at first colourless, but by exposure to air it speedily becomes brown and opaque, from the conversion of the white soluble prothydrate into the brown insoluble sesquihydrate. Manganese compounds are very frequently contaminated with iron; hence, when excess of ammonia produces a brown precipitate, it is often worth while to pour the mixture on to a filter, and to test the ammoniacal filtrate with sulphide of ammonium; whereupon manganese, if present, is precipitated as a flesh-coloured sulphide. Manganese is best recognised before the blowpipe by fusing some carbonate of sodium, to which a little nitre may be added, upon platinum foil,

and then introducing a minute portion of the manganese compound, whereby a bright green fusible mass of manganate of sodium Na_2MnO_4 is produced.

(40.) δ . The potash solution of chromium is of a green colour, and when boiled for a longer or shorter time, according to the relative proportions of potash and chrome present, deposits the whole of its chrome in the form of a green precipitate. Chrome compounds are best recognised by the emerald green colour they impart to the borax bead; and by the bright yellow mass of chromate of alkali metal which they yield when ignited on platinum foil, with carbonate of sodium and nitre, before the blowpipe.

ϵ . The potash solution of zinc, when containing a considerable excess of potash, is not affected even by prolonged ebullition; but when containing only a slight excess, it soon deposits an opaque white precipitate. On adding a few drops of *sulphide of ammonium* to potash solution of zinc, sulphide of zinc is formed, which being, unlike the hydrate of zinc, insoluble in potash, is thrown down as a white precipitate, frequently having a greyish tinge from the accidental presence of a trace of lead in the potash used as the reagent.

Zinc compounds when strongly heated leave a fixed infusible residue, which when cold is quite white, but when hot has a deep greenish-yellow tint. Moistened with nitrate of cobalt solution and re-ignited, it acquires a pure permanent chrome-green colour.

ζ . The potash solution of alumina is unaffected by boiling, and also by the addition of a few drops of sulphide of ammonium, unless indeed the potash contains an appreciable quantity of lead, when a black precipitate of sulphide of lead is formed.

Ammonia or carbonate of ammonium gives a similar white precipitate with zincous and aluminous salts, but the precipitate produced with the former is freely soluble, while that produced with the latter is insoluble in excess of the reagent. Hence the addition of *chloride of ammonium* to potash solution of zinc

produces no precipitate ; but when added in sufficient quantity to potash solution of alumina, it throws down a white gelatinous precipitate of hydrate of aluminum, owing to a substitution of free ammonia for free potash, thus :



Compounds of aluminum, when strongly heated in the blowpipe flame, leave a white, infusible, highly incandescent residue, which, when moistened with solution of nitrate of cobalt, and reignited, assumes a fine permanent blue colour.

§ V.—EXAMINATION FOR BASES OF GROUP III.

(41.) This group includes the bases which are not precipitated either by sulphuretted hydrogen, or by sulphide of ammonium. A solution of the salt in water or acid may be at once examined, according to the directions of Table III. Or the solution, which has been successively treated with sulphuretted hydrogen and sulphide of ammonium without the production of a precipitate, may be tested for any of the metals of this group, except potassium, sodium, and ammonium. If the original substance required an acid to effect its solution, its base cannot be any one of the last three, inasmuch as all simple salts of potassium, sodium, and ammonium are soluble in hot water.

TABLE III.

(42.) Examination of a solution containing some one member of the third group of bases; namely, BARIUM, STRONTIUM, CALCIUM, MAGNESIUM, POTASSIUM, SODIUM, or AMMONIUM. The sulphides or sulphhydrates of these metals being soluble in water, the solutions of their salts are not precipitated either by sulphuretted hydrogen, or by sulphide or sulphhydrate of ammonium.

a. Add *Chloride and Carbonate of Ammonium* to the original solution, and warm gently.

Precipitated

BARIUM

STRONTIUM

CALCIUM.

Dissolved

MAGNESIUM

POTASSIUM

SODIUM

AMMONIUM.

b. Add *Sulphate of Potassium* or dilute *Sulphuric Acid* to the original solution.

γ. Add *Phosphate of Ammonium* to the above solution already containing the ammoniacal salts, and stir.

Precipitated

BARIUM (*quickly*)

STRONTIUM (*slowly*).

Dissolved

CALCIUM.

Add *Chromate of Potassium* to the original solution.

Precipitated

MAGNESIUM.

Dissolved

POTASSIUM

SODIUM

AMMONIUM.

δ. Add to the original solution *Tartaric Acid* in excess, and stir.

Precipitated

BARIUM.

Dissolved

STRONTIUM.

Precipitated

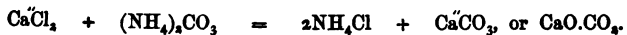
POTASSIUM.

Dissolved

SODIUM.

AMMONIUM ?

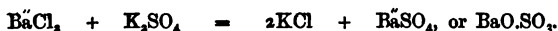
(43.) *a. Carbonate of ammonium* produces in solutions containing barium, strontium, or calcium, a white precipitate, the carbonates of these three metals being insoluble :



When carbonate of ammonium is added to an acid solution of the compound under examination, care must be taken to add a quantity more than sufficient to neutralise the acid, or, in other words, enough to render the solution alkaline. Inasmuch as the carbonic gas evolved on adding carbonate of ammonium to an acid liquid retains the carbonates of the alkaline earth-metals in solution, it must be expelled by gentle warming. Carbonate of magnesium also is insoluble in water, though readily soluble in solutions containing chloride of ammonium or other ammoniacal salt. Hence the necessity of adding chloride as well as carbonate of ammonium to the aqueous solution of the substance under examination, in order to prevent a precipitate of carbonate of magnesium being formed, and confounded with one of the other precipitable carbonates. But when the substance has been dissolved in an acid, the carbonate of ammonium expended in neutralising the acid forms a sufficient quantity of neutral ammoniacal salt to prevent the precipitation of carbonate of magnesium ; and consequently the separate addition of chloride of ammonium is rendered unnecessary. The addition of chloride of ammonium may also be dispensed with, when the carbonate is added to a portion of the solution with which a negative result has been obtained by successive treatment with sulphuretted hydrogen and sulphide of ammonium. Carbonate of ammonium also precipitates most of the metals belonging to the first and second groups, so that it can only be depended upon to indicate the presence of the alkaline earth-metals, when the absence of all other precipitable metals has been previously ascertained by sulphuretted hydrogen and sulphide of ammonium respectively.

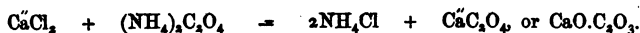
β. Sulphate of calcium is very slightly soluble in water, sulphate of strontium still less so, and sulphate of barium

quite insoluble. Hence sulphuric acid and most soluble sulphates give precipitates in the solutions of all three metals. But owing to the sparing solubility of *sulphate of potassium* in water, its solution does not usually precipitate the salts of calcium unless very concentrated, though it is readily capable of precipitating those of strontium and barium :



Moreover, a solution of sulphate of calcium will not precipitate salts of calcium under any circumstances, but will nevertheless precipitate those of strontium and barium, the former slowly, the latter immediately.

Oxalate of ammonium is the reagent usually employed to demonstrate the presence of calcium, with neutral or alkaline solutions of which it gives a white precipitate of oxalate of calcium, soluble in nitric and hydrochloric, but insoluble in acetic and oxalic acids :



But oxalate of ammonium produces precisely similar precipitates in neutral salts of strontium and barium, the oxalates of these two metals being also insoluble in water. That of barium, however, is soluble in excess of warm oxalic acid. Moreover, free oxalic acid precipitates neutral solutions of calcium and strontium, but not of barium, unless very concentrated.

In solutions acidified with acetic acid, *chromate of potassium* has no action upon strontium and calcium salts, but throws down from barium salts a yellow precipitate of chromate of barium, soluble in the nitric and hydrochloric acids :



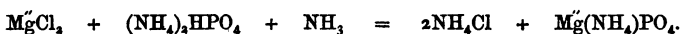
Hydrofluosilicic acid also serves to distinguish barium salts from those of strontium and calcium, with the first of which it alone produces a precipitate, the fluosilicate of barium being alone

insoluble. This test is unaffected by the presence of nitric and hydrochloric acids :



Most barium salts, especially when moistened with hydrochloric acid, impart an apple-green colour to the blowpipe flame; strontium salts a marked crimson; and calcium salts an orange-red.

(44.) *γ. Phosphate of ammonium, or of sodium*, gives with magnesian solutions, containing carbonate of ammonium or free ammonia, a white crystalline precipitate of phosphate of magnesium and ammonium, frequently known as triple phosphate :



This precipitate does not usually form in warm solutions, and is frequently produced only after brisk stirring, or rubbing the inside of the tube with a glass rod. It must be remembered that phosphate of ammonium gives precipitates with salts of barium, strontium, and calcium, and of most metals belonging to the first and second groups, so that it can only be relied upon as a test for magnesium when the absence of all other metals precipitable by it has been first ascertained.

δ. Tartaric acid gives with neutral solutions of potassium-salts a white crystalline precipitate of acid tartrate of potassium or cream of tartar :



The precipitate does not usually appear in a warm liquid. It is produced very slowly in solutions of the more sparingly soluble, and in dilute solutions of the more freely soluble potassium-salts. From such solutions it is best obtained by cooling the mixture under a tap, diluting it with a little alcohol, and stirring it briskly with a glass rod, which should be rubbed

against the inside of the tube. In testing alkaline solutions for the presence of potassium, care must be taken to add an excess of tartaric acid. The precipitate of cream of tartar is not formed in solutions of potassium-salts which have a marked acid reaction from the presence either of oxalic acid, or of one of the strong mineral acids. But on neutralising such solutions with caustic soda, or carbonate of sodium, and then adding tartaric acid, the characteristic precipitate is readily obtained.

Perchloride of platinum produces also in solutions of potassium-salts, which should first be acidulated with hydrochloric acid, a yellow crystalline precipitate of platino-chloride of potassium 2KCl.PtCl_4 , the deposition of which is facilitated by stirring, and by the addition of alcohol.

Solutions of ammonium-salts, when moderately concentrated, resemble potassium-salts in their behaviour with tartaric acid, yielding therewith a crystalline precipitate of acid tartrate of ammonium $(\text{NH}_4)_2\text{H}_2\text{C}_4\text{O}_6$. Hence ammonia cannot be used to neutralise those acid solutions which have to be tested with tartaric acid for the presence of potassium. Moreover, moderately concentrated solutions of ammonium-salts give with dichloride of platinum a yellow crystalline precipitate of platino-chloride of ammonium $2\text{NH}_4\text{Cl.PtCl}_4$, resembling the similar compound of potassium. Dilute solutions of ammoniacal salts behave like solutions of sodium-salts in not furnishing any precipitate with either tartaric acid or dichloride of platinum. But ammonium-salts may be readily distinguished from those of potassium and sodium by their partial or complete volatility, and by their solutions evolving ammonia when boiled with potash or lime:



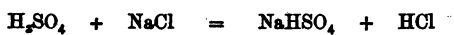
Sodium-salts impart an intense yellow colouration to flame, and potassium-salts a marked violet.

§ VI.—EXAMINATION FOR ACIDS.

(45.) The student should always ascertain the nature of the metallic or basic constituent of his substance, before proceeding to search for its acid or chloroid constituent. In this search he will be greatly assisted by a knowledge of the special characters of particular salts, and of the general characters of various classes of salts. Knowing, for instance, that sulphate of barium is insoluble in all menstrua, he need not test a soluble barium salt for sulphuric acid. On the other hand, a salt insoluble in water is not likely to be a nitrate or chlorate. The following classes of salts are, as a rule, soluble in water:—Nitrates, excepting a few superbasic salts; acetates, except acetate of silver, which is only sparingly soluble; chlorides, except chloride of silver, which is insoluble in boiling nitric acid, and mercurous chloride, or calomel, which dissolves in boiling nitric acid with conversion into a mercuric salt: chloride of lead is moderately soluble in boiling, though very sparingly soluble in cold water. Sulphates, except those of barium, strontium, and lead, which last is soluble in boiling hydrochloric acid: the sulphates of silver and calcium also are only sparingly soluble. The following classes of salts are, as a rule, insoluble in water:—Oxides or hydrates, and sulphides or sulphydrates, except those of the alkali- and alkaline earth- metals, and the sulphydrate of magnesium; also carbonates, phosphates, and oxalates in general, except those of the alkali metals, and many hyperacid salts.

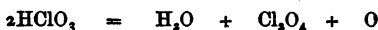
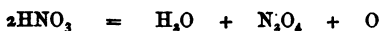
The presence of any particular acid is indicated more or less certainly by the behaviour of the substance when gently heated in the solid state with three or four times its bulk of strong *sulphuric acid*, and when under examination for the detection of its metallic constituent.

The most usual decomposition which takes place between sulphuric acid and the salt of a more volatile or feeble acid consists in an exchange of the hydrogen of the sulphuric acid for the metal of the salt, thus :



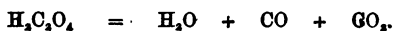
Reactions similar to the above take place with the salts of nearly all acids, provided the sulphuric acid employed is not too strong, and the heat not too great; but with ordinary oil of vitriol, especially at a somewhat high temperature, the liberated acids themselves often undergo a partial or complete decomposition.

The tartaric and citric acids, for instance, are destroyed with more or less charring. The nitric and chloric acids yield the peroxides of nitrogen and of chlorine respectively, thus:

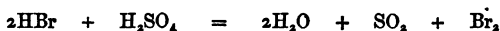


except that, in the former case, the decomposition is but slight, while in the latter, the separated oxygen converts another portion of the chloric into perchloric acid.

Oxalic acid breaks up into water, carbonous oxide, and carbonic anhydride, thus:



The hydrobromic, hydroiodic, and sulphydric acids are decomposed with liberation of bromine, iodine, and sulphur, respectively, thus:



Hydrocyanic acid yields acid-sulphate of ammonia and carbonous oxide, thus:



(46.) In the following table the principal acids are classified according to the behaviour of their salts when under examination for bases, and when warmed with strong sulphuric acid:

TABLE IV.—PRELIMINARY TESTING FOR ACIDS.

α. Indicated during previous examination.	β. Corroborated by sulphuric acid.	γ. Also react with sulphuric acid.	δ. No obvious effect with sulphuric acid.
ARSENATES } React CHROMATES } with H_2S	NITRATES. Brown acid vapours with copper	CHLORIDES. Pungent acid fumes. With MnO_2 evolve chlorine	BORATES. Give green flame with alcohol
NITRATES } Defla- CHLORATES } grate	CHLORATES. Brown ing and crackling detonation	BROMIDES. Brown fumes, which bleach litmus, and turn starch yellow.	OXALATES } Efferv- CYANIDES } vesce?
TARTRATES } Char CITRATES }	TARTRATES } Char CITRATES }	IODIDES. Violet vapours, which turn starch paper purple.	PHOSPHATES } ARSENATES } SULPHATES } SILICATES } OXIDES }
CARBONATES } Efferv- SULPHIDES } vesce SULPHITES, &c. } with PEROXIDES } HCl	CARBONATES } Efferv- SULPHIDES } vesce SULPHITES }	FLUORIDES. Pungent acid fumes, which etch glass	
PERSULPHIDES } SILICATES } Give precipi- BORATES } tates with BENZOATES } HCl		ACETATES. Acid vapour. With alcohol yield acetic ether	

(47.) α. Deflagration of the substance, when heated upon charcoal before the blowpipe, shows the probable presence of a nitrate or chlorate.

Charring of the substance, when heated upon platinum foil or charcoal, usually indicates the presence of a tartrate or citrate.

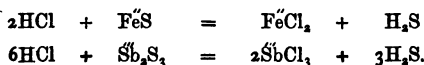
Effervescence from the substance, when its solution is being effected in hydrochloric acid, occurs with several classes of salts.

With nearly all carbonates there is evolution of carbonic anhydride, thus :



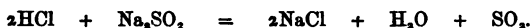
The gas is free from any marked smell, and renders lime water milky (*vide par.* 84).

With many sulphides there is evolution of sulphydric acid or sulphuretted hydrogen, thus :



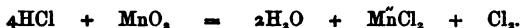
The gas smells most offensively, and turns lead paper of a black or brown colour.

With sulphites and hyposulphites (*vide par.* 94) there is evolution of sulphurous anhydride, thus :



The gas has the peculiar suffocating smell of burning sulphur, and produces a purple discolouration on starch paper moistened with iodic acid solution.

With peroxides there is evolution of chlorine, thus :



The gas, which has a greenish colour and characteristic irritating smell, produces a purple colouration on starch paper moistened with iodide of potassium.

Similar effects are observable on acidifying with hydrochloric acid the strong aqueous solutions of soluble carbonates, sulphides or sulphydrates, sulphites, and hyposulphites ; but the evolution of chlorine under these circumstances would probably be due to the presence of a hypochlorite. Moreover, most simple cyanides, and the strong aqueous solutions of such of them as are soluble, effervesce when warmed with hydrochloric acid, from an evolution of hydrocyanic acid gas, known by its characteristic smell, and by its behaviour with sulphide of ammonium, &c. (*vide par.* 99).



The aqueous solutions of certain salts when acidified with

hydrochloric acid yield precipitates due, not to the metals silver, lead, and mercury, but to the acids of the respective salts.

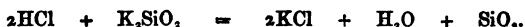
With persulphides there is formed a yellowish white turbidity from liberated sulphur, always accompanied, however, by an evolution of sulphuretted hydrogen :



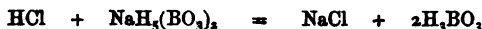
With hyposulphites, there is gradually produced a more decidedly yellow precipitate of sulphur, and an evolution of sulphurous anhydride, thus :



With soluble silicates there is produced a gelatinous precipitate of silica, which separates out more completely, and becomes gritty on evaporation :



Borates and benzoates, if sufficiently concentrated, yield crystalline precipitates of boric and benzoic acids respectively, which dissolve in boiling water, and crystallise out again on cooling :

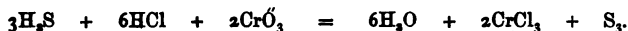


In testing for bases of the first group, by treating the acidified solution of the substance with sulphuretted hydrogen, strong evidence of the presence of arsenates or chromates may also be afforded. Acidified solutions of arsenates slowly yield a yellow precipitate of mixed trisulphide of arsenic and sulphur, thus :

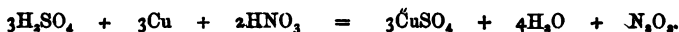


Acidified solutions of chromates yield a yellowish-white deposit of sulphur, while the colour of the liquid changes from

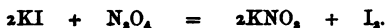
orange-red to green, owing to the reduction of the chromic acid to the state of a chrome-salt, thus :



(48.) β . Nitrates, when heated with sulphuric acid, give off nitric acid vapours, sometimes having a brownish tint from the presence of peroxide of nitrogen. On the addition of a few copper turnings the brown colour becomes very marked, from the evolution of colourless nitric oxide N_2O , and its immediate conversion into brown peroxide of nitrogen N_2O_4 , by an absorption of atmospheric oxygen :



The vapour, whether before or after the addition of copper, has a characteristic nitrous smell, reddens litmus paper, and gives a purple colouration to starch paper moistened with iodide of potassium :



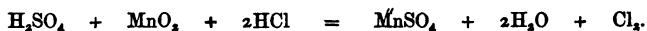
Chlorates react somewhat violently with sulphuric acid, producing an immediate browning, and, either at once or on the slightest application of heat, a crackling, or even loud detonation, from the breaking up of the peroxide of chlorine first liberated.

Tartrates and citrates undergo more or less charring, the tartrates becoming thoroughly blackened, the citrates merely browned.

Carbonates, sulphites, and sulphides effervesce with sulphuric acid as with hydrochloric acid ; except that in the last case some of the liberated sulphydric acid reacts with the excess of sulphuric acid to form sulphur, water, and sulphurous anhydride, as already described (vide par. 45).

γ . Most chlorides when warmed with sulphuric acid evolve hydrochloric acid gas, which reddens litmus paper, and has a well characterised pungent smell. On the addition of a little peroxide

of manganese, chlorine is evolved, recognisable by its peculiar irritant smell :—



It bleaches litmus, and produces a purpling of starch paper moistened with iodide of potassium solution.

Bromides evolve a mixture of hydrobromic acid with bromine vapour, the latter recognisable by its brown colour, irritant smell, and power of bleaching litmus.

Iodides yield iodine vapour, recognisable by its violet colour, and by its rendering starch paper purple.

Fluorides give off pungent fumes of hydrofluoric acid, which redden litmus and etch glass (*vide par.* 106).

Acetates give off acetic acid vapours, which redden litmus. On the addition of a little alcohol the original sour smell of the acid is changed into the fruity smell of acetic ether.

δ. Borates do not react visibly with sulphuric acid, but on adding alcohol, and then setting fire to the mixture in a capsule, the flame presents a marked green colour.

Oxalates and cyanides are decomposed by heated sulphuric acid with liberation of carbonous oxide gas, accompanied in the case of the former salts with carbonic anhydride; but as the evolution of gas is liable to be overlooked, and the gas itself does not present any striking property, the oxalates and cyanides are here included among the salts with which sulphuric acid produces no obvious effect.

Phosphates, arsenates, sulphates, silicates, and oxides do not react visibly with sulphuric acid; except that with some peroxides there is evolution of oxygen, transformable into that of chlorine on the addition of chloride of sodium or hydrochloric acid.

(49.) The various liquid tests for the acids should be applied by preference to an aqueous solution of the original substance. But if insoluble in water, and consequently neither a nitrate nor a chloride, it may be dissolved in dilute nitric or hydrochloric acid, and the tests applied to the solution so formed.

The presence of certain basic metals interferes occasionally with the described reactions of several of the acids. Thus solution of hydrochloric acid or any chloride, when tested with nitrate of silver, gives a white precipitate of chloride of silver, said to be readily soluble in ammonia. But on adding nitrate of silver to solution of mercuric chloride, and treating the resultant white precipitate with ammonia, there is no obvious solution produced, because, although the precipitated chloride of silver does actually and completely dissolve, a white insoluble mercurammonium compound is simultaneously thrown down from the mutual reaction of the ammonia and mercuric salt.

Bearing in mind, however, the interference likely to be produced by the presence of the particular metal previously detected, the tests for sulphuric, hydrochloric, and nitric acids can for the most part be satisfactorily applied to any of their salts.

But in testing for phosphoric, oxalic, and tartaric acids more particularly, and for other acids when any difficulty occurs, it is important that the base of the salt should be one of the alkali metals, potassium, or sodium, or ammonium.

The bases of the first group may be got rid of by saturating the solution with sulphuretted hydrogen gas, filtering, evaporating the filtrate until it ceases to smell, and neutralising it with carbonate of sodium or potassium.

The bases of the second group, together with barium, strontium, calcium, and magnesium, may be removed by adding to the acidulous, or occasionally to the aqueous, solution of the substance an excess of carbonate of sodium or potassium, boiling for some time, and filtering. The filtrate can afterwards be neutralised with nitric or hydrochloric acid, which may be conveniently added drop by drop from a pipette.

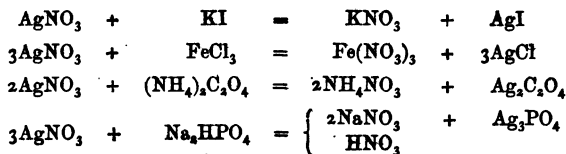
(50.) The following abridged table shows the action of some general reagents upon the dissolved salts of the principal acids. By its aid and that of the preceding table, the student will rarely have much difficulty in quickly discovering the acid constituent of his substance, although, indeed, the course of examination is not so systematic as that for the bases. A more complete table

for the detection of the acids would include the reactions of a few organic salts of comparatively rare occurrence, such as the formates, succinates, citrates, meconates, gallates, tannates, and ferrid-cyanides; of a few mineral salts of similarly rare occurrence, namely, the iodates, seleniates, and silico-fluorides; of a few mineral salts rarely met with in the soluble form, namely, silicates and fluorides; and of the previously detected arsenates. It would also mention the several precipitates given by nitrate of silver and nitrate of barium respectively, which disappear on acidification, and have, as a rule, but little practical interest.

TABLE V.—COURSE FOR DETECTION OF THE ACIDS.

a. NITRATE OF SILVER	β. NITRATE OF BARIUM	γ. CHLORIDE OF CALCIUM	ε. PERCHLORIDE OF IRON
From acid solutions	From acid solutions	From acetic acid sols.	From acid solutions.
CHLORIDES } CYANIDES } White BROMIDES }	SULPHATES. White	OXALATES. White	FERROCYANIDES. D. blue pp.
IODIDES. Yellow	From non-acid sols.	From neutral sols.	SULPHOCYANIDES. D. red colour.
SULPHIDES. Black	CHROMATES. Yellow	TARTRATES. White	From neutral sols.
	CARBONATES } SULPHITES } White &c., &c.	δ. SULPHATE OF MAGNESIUM	BORATES } L. brown BENZOATES } pp.
		From ammon. sols.	ACETATES. D. red colour
		PHOSPHATES. White	

(51.) a. *Nitrate of silver* causes precipitates in the solutions of very many classes of salts, the majority of silver salts being more or less insoluble in water. The reactions consist in an exchange of the silver of the nitrate of silver for the metal or quasi-metal of the dissolved salt under examination, as illustrated by the following examples :—



All the precipitates produced by nitrate of silver disappear upon the addition of nitric acid, or do not form in presence of free nitric acid, except the chloride, cyanide, bromide, iodide, and sulphide.

With chlorides the silver precipitate is white, becoming slate-coloured on exposure to light, and soluble in ammonia before discolouration. Upon heating the original substance with peroxide of manganese and sulphuric acid, chlorine gas is evolved.

With simple* cyanides the silver precipitate is white, soluble in ammonia, and in boiling concentrated nitric acid: a portion of the precipitate washed by decantation may be treated with yellow sulphide of ammonium, whereby sulphide of silver and sulphocyanate of ammonium are produced, which last strikes a deep red colour on the addition of a ferric salt. An excess of sulphide of ammonium must be avoided, or afterwards got rid of by evaporating to dryness.

With bromides the silver precipitate is white, and with difficulty soluble in ammonia. Bromine is liberated from the original salt, when treated with sulphuric acid, and from the solution of the salt when treated with a few drops of nitro-hydrochloric acid or chlorine-water, whereby the liquid assumes a red-brown colour rendered more evident upon the addition of a little ether or chloroform, which dissolves out the bromine to form a deep brown stratum.

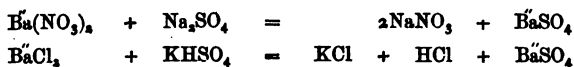
With iodides the silver precipitate has a pale yellow colour, and is insoluble in, but turned white by, ammonia. Iodine is liberated from the salt when treated with sulphuric acid, and from the solution of the salt when treated with a few drops of yellow nitric or nitro-hydrochloric acid or chlorine-water, its presence being manifested by the purple colour it produces on starched paper, or with dilute starch paste added to the liquid, or by the pink or crimson colour it imparts to chloroform.

* Ferrocyanide and sulphocyanate of silver occur as white precipitates, and the ferridcyanide as a brown precipitate, all unaffected by nitric acid; but soluble ferrocyanides, ferridcyanides, and sulphocyanates are recognised immediately by their behaviour with iron solutions.

With sulphides the silver precipitate is black, and insoluble in ammonia, but soluble in boiling nitric acid. Nitro-prusside of sodium added to an alkaline sulphide produces a deep purple colouration.

Of the silver precipitates which disappear on acidification with nitric acid, the hydrate is brown, the chromate dark red, the arsenate brick-red, the phosphate bright yellow, though sometimes white, the carbonate pale yellow, and the rest white. The oxalate is insoluble in acetic acid; the acetate is thrown down from concentrated solutions only; the tartrates, formates, and sulphites are reduced to the metallic state on boiling; while the borates, benzoates, and citrates do not exhibit any characteristic property.

(52.) β . *Nitrate or chloride of barium* precipitates the solutions of many classes of salts, most barium salts being insoluble or sparingly soluble in water. The reaction consists in an exchange of the barium of the nitrate or chloride of barium for the metal or quasi-metal of the salt under examination, thus:



With sulphates the barium precipitate is white, and, if in any quantity, opaque. It does not disappear upon the addition of nitric or hydrochloric acid, but is nevertheless slightly soluble in concentrated nitric acid. The seleniate and silicofluoride of barium also occur as white precipitates unaffected by acidification, (vide par. 95). Of the barium precipitates which dissolve in nitric or hydrochloric acid, the chromate is yellow and the remainder are white. The carbonate and sulphite dissolve with effervescence. The arsenate, borate, and tartrate do not form in the presence of ammoniacal solutions, and, when once thrown down, disappear more or less readily on the addition of chloride of ammonium. The oxalate and phosphate exhibit no characteristic properties.

(53.) γ . With a few exceptions, *chloride of calcium* causes precipitates with the several classes of salts which are precipitated by chloride or nitrate of barium; but while sulphate of barium

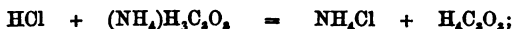
is much more insoluble in water and acids than sulphate of calcium, oxalate of calcium is more insoluble in water, and especially in acetic acid, than oxalate of barium.

With solutions of oxalates in which there is no free mineral acid, chloride or sulphate of calcium produces an opaque white precipitate of oxalate of calcium, the deposition of which, when in small quantity, may be facilitated by stirring:



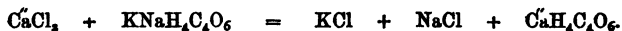
The precipitate is insoluble in ammonia and in acetic acid, but soluble in dilute nitric or hydrochloric acid. Its solubility in mineral acids distinguishes it from sulphate of calcium, which is occasionally thrown down by chloride of calcium from strong solutions of sulphates, as does also its property of effervescing with sulphuric acid and peroxide of manganese; while its insolubility in acetic acid distinguishes it from the phosphate, tartrate, carbonate, and other salts of calcium insoluble in water.

The solution to be tested for oxalic acid with chloride of calcium may be alkaline, or neutral, or acid from acetic acid only. If an alkaline carbonate is present from its employment in preparing the solution, it should be neutralised with acetic or hydrochloric acid. If an excess of mineral acid is present, it may either be neutralised with ammonia, or acetate of ammonia may be added, whereby the free mineral acid is replaced by free acetic acid, thus:



but in practice it is better to add the ammonia and acetic acid separately.

In neutral solutions of tartrates, chloride of calcium gives a white precipitate of tartrate of calcium, soluble even in acetic acid:



On digesting the precipitate with potash, filtering, and boiling the filtrate, a white turbidity is produced, which disappears on

cooling. This behaviour with potash distinguishes the tartrate from the phosphate and other insoluble salts of calcium. Tartrate of calcium is not thrown down in the presence of much ammoniacal salt.

(54.) *δ*. Although phosphate solutions are precipitated alike by nitrate of silver, nitrate of barium, chloride of calcium, sulphate of magnesium, and perchloride of iron, it is the magnesian precipitate more especially which is characteristic of phosphoric acid. From a neutral or alkaline phosphate solution, a mixture of *sulphate of magnesium*, chloride of ammonium and ammonia throws down a white crystalline precipitate of phosphate of ammonium and magnesium, thus :



In common with all magnesian precipitates, it is readily soluble in acids ; but, unlike any other magnesian salt, except the arsenate, it is insoluble in ammonia and ammoniacal salts. The yellow precipitate given with nitrate of silver is a useful confirmatory test, as is also the reaction with molybdate of ammonium (*vide par.* 100), although this last is common to the phosphoric and arsenic acids.

(55.) *ε*. *Perchloride of iron*, which, for most purposes, should be free from any excess of acid, causes precipitates or characteristic alterations of colour in the solutions of very many classes of salts, among which may be mentioned the following :—

With ferrocyanides a dark blue precipitate of Prussian blue, $\text{Fe}_3\text{Cy}_{18}$ or $3\text{Fe}''\text{Cy}_{2.4}\text{Fe}'''\text{Cy}_3$, is thrown down ; while with sulphocyanates a dark red colour, due to ferric sulphocyanate $\text{Fe}''(\text{CyS})_3$, is developed, neither of which results is affected by the addition of hydrochloric acid. With neutral borates, benzoates, or phosphates, there is produced a pale brown precipitate of borate, benzoate, or phosphate of iron, which disappears on the addition of hydrochloric acid. The phosphate, however, $\text{Fe}'''\text{PO}_4$, is scarcely affected by acetic acid, save in presence of a large excess of iron.

With neutral acetates and sulphites, there is produced a

dark reddish-brown colouration, which disappears on the addition of hydrochloric acid, or on boiling: in the latter case with formation of a red-brown turbidity.

Nitrates and chlorates are unaffected by any of the general reagents mentioned in the table.

(56.) Although nearly all of the acids react more or less characteristically with several reagents as above described, the following list of acids or salts, with the tests by which their presence is more particularly indicated or established, may prove useful to the student:—

CHROMATES	Reaction with sulphydric acid.
NITRATES	} Deflagration on charcoal, and reaction with sulphuric acid.
CHLORATES	
CARBONATES,	} Effervescence with acids.
SULPHIDES	
SULPHITES	
SULPHATES	Precipitation by chloride of barium.
CHLORIDES	} Precipitation by nitrate of silver.
BROMIDES	
IODIDES	
CYANIDES	
PHOSPHATES	Precipitation by sulphate of magnesium.
OXALATES	} Precipitation by chloride of calcium.
TARTRATES	
ACETATES	} Reaction with perchloride of iron.
BENZOATES	
BORATES	} Special tests.
FLUORIDES	
SILICATES	

In the paragraphs, 89 to 107, describing the several reactions of the individual acids, the order of the above list is adopted.

§ VII.—SPECIAL SUBSTANCES.

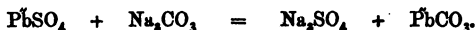
(57.) It being scarcely possible to give general rules that shall apply to every particular case, there yet remain for consideration certain compounds, the detection of which according to the directions of the preceding tables is impracticable, or difficult, or liable to fallacy. There are some few substances, for instance, which are insoluble in all ordinary menstrua, and which must consequently be submitted to special modes of treatment, so that they may be transformed into new and soluble combinations. Again, there are those compounds of barium, strontium, calcium, and magnesium, which are precipitated by sulphide of ammonium along with the members of the second group, and which are known by the laboratory-name of earthy salts. There are also the oxides, or compounds without an acid, and the acids or compounds without a base other than hydrogen, together with two or three miscellaneous bodies. A few remarks, moreover, are appended on the examination of substances in the dissolved or liquid condition, and on the solubility of the heavy metals in alkaline solutions.

(58.) INSOLUBLE COMPOUNDS.

Many substances are known which do not dissolve in water or in any ordinary acid, or in any combination of acids, even with the aid of heat. Among bodies of this class, the following are most likely to come under the notice of the student: namely, peroxide of tin; peroxide of antimony; chloride, bromide, and iodide of silver; the sulphates of barium, strontium, and perhaps lead; chromic oxide; alumina and some aluminates; with silica and some silicates. Several of these bodies may be met with in the soluble as well as the insoluble form, but the silver compounds and earthy sulphates are always alike insoluble. Sulphate of strontium, however, though practically insoluble, dissolves sufficiently in water to form a solution in which the presence

of sulphuric acid can be detected by the addition of a barium-salt. Again, sulphate of lead dissolves, though not very readily or to any great extent, in boiling hydrochloric acid, forming a solution from which chloride of lead crystallises out on cooling. For analytical purposes these more or less insoluble bodies may be classified into compounds of heavy metals easily recognisable before the blowpipe (α), and earthy compounds for whose satisfactory identification a further examination is required (β).

α . The insoluble compounds of tin, antimony, silver, and lead, when mixed with cyanide flux and heated on charcoal before the blowpipe, yield beads of metal distinguishable from one another by their appearance, texture, behaviour with acids, and freedom from, or association with, incrustations of definite character (vide par. 26, δ). The sulphuric acid of the insoluble lead compound may be recognised by boiling it with a solution of carbonate of sodium or of potassium, filtering, and testing the clear filtrate, with nitric acid and nitrate of barium, for the presence of any alkaline sulphate resulting from the decomposition of the original sulphate of lead, thus:



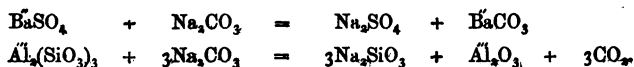
The chlorine, bromine, or iodine of the insoluble silver compound may be detected by fusing the compound with a mixture of the carbonates of sodium and potassium in a small porcelain capsule or iron spoon, and treating the resulting saline mass with sulphuric acid and peroxide of manganese, or the solution of the mass with nitric acid and nitrate of silver. By fusing the chloride, bromide, or iodide of silver with carbonated alkali, the halogen is transferred from the silver to the alkali metal, thus:—



Chromic oxide is known by its dull green colour, by the bright green colour it imparts to the borax bead, and by the yellow mass of alkaline chromate into which it is converted by fusion with a little nitre and carbonate of sodium. This mass

may be dissolved in water, and to the yellow solution so formed the ordinary tests for chromic acid applied.

β. The remaining insoluble and non-reducible compounds may be recognised by the following process. The substance in the state of a very fine powder, and mixed with three or four times its weight of the carbonates of sodium and potassium, is fused for some time on a platinum capsule or on platinum foil, whereby a decomposition of the kind represented below is effected :



By boiling the fused mass in water and filtering, a clear solution is obtained containing a sulphate or silicate, and perhaps some aluminate of sodium; while an undissolved residue is left upon the filter consisting of alumina, or of carbonate of barium or strontium, or possibly of the carbonate or oxide of some other basylous metal, originally existing in the form of a silicate or aluminate. This residue, after being washed with water, is dissolved in dilute hydrochloric acid, and the resulting solution tested by the usual reagents, for aluminum, barium, strontium, and the basylous metals generally.

The clear filtrate is acidified with hydrochloric acid, and a portion of it, refiltered if necessary from any deposit caused by the acidification, tested for sulphuric acid by means of nitrate or chloride of barium. The remainder, whether or not turbid from the acidification, must be evaporated to dryness, gently heated, and again acted upon with hydrochloric acid and water. Any undissolved gritty residue will consist of silica, which may be further identified by fusion with a bead of carbonate of sodium before the blowpipe; while any alumina contained in the hydrochloric acid solution will be thrown down on neutralisation with ammonia.

(59.) EARTHY SALTS.

These compounds being insoluble in water are precipitated unchanged from their acidulous solutions upon the addition thereto of any alkaline hydrate, carbonate, or sulphide; whence their occurrence among members of the second group of bases. Perphosphate of iron, which resembles the earthy salts in the behaviour of its acidulous solution with alkaline hydrates and carbonates, may be conveniently associated with them, thus :

α. Iron	Phosphate
Barium	}	Phosphate
β. Strontium		Oxalate
Calcium		Fluoride
γ. Magnesium	Phosphate

The identification of the different compounds when separated from one another is easily effected by the under-mentioned methods; while the more difficult process for the detection of iron, calcium, magnesium, and phosphoric acid in presence of one another is described in Chapter IV.

α. In the original acid solution, iron may be detected by sulphocyanate or ferridecyanide of potassium; and phosphoric acid by molybdate of ammonium and nitric acid. After any excess of mineral acid, more than sufficient to keep the substance in solution, has been neutralised with ammonia, the addition of acetate of ammonium produces a buff-coloured precipitate of phosphate of iron, unless, indeed, the ratio of phosphoric acid to iron is very small.

β. Barium and strontium may be detected in the original solution by means of sulphate of potassium, and be distinguished by the different behaviour of their salts, with hydrofluosilicic acid, or before the blowpipe. Calcium may also be detected in the original solution by the addition thereto of sulphate of potassium and proof spirit. The precipitated sulphate of calcium may be collected on a filter, washed with proof spirit, dissolved in water, and the resulting solution tested with oxalate of ammonium.

Phosphoric acid may be detected in the original solution by molybdate of ammonium and nitric acid.

Oxalic acid is recognisable by the effervescence produced on adding dilute sulphuric acid and peroxide of manganese to the original substance. Or it may be detected by boiling the original acid solution with excess of carbonate of sodium, filtering, rendering the filtrate slightly acid with acetic acid, and then adding chloride of calcium, when a white precipitate of oxalate of calcium will be produced. Moreover, like phosphate of iron, oxalate of calcium is thrown down from its original solution in hydrochloric acid by addition of acetate of ammonium.

Fluorine may be detected by treating the original substance with sulphuric acid, or with sulphuric acid and silica (*vide par.* 106).

γ. Magnesium in any form is not precipitated from acid solutions by sulphate of potassium and proof spirit, or by acetate of ammonium; but, in presence of phosphoric acid, is thrown down from its acid solution by excess of ammonia as a white crystalline deposit of triple phosphate, which may be examined microscopically, and if necessary be dissolved in acetic or hydrochloric acid, so as to form a solution that may be submitted to further examination.

(60.) OXIDES AND SULPHIDES.

The oxides are generally recognised by their physical properties, and by their not answering to the tests for any of the acids. Peroxides when boiled with hydrochloric acid, or with sulphuric acid and common salt, give off chlorine gas. The soluble hydrates or hydrated oxides are known by their alkalinity to test paper; by their effervescing not at all, or but very slightly, upon the addition of an acid; and by their giving a brown precipitate with solution of nitrate of silver. The solution of a hydrate is, moreover, distinguishable from that of a carbonate by the circumstance that an admixture with it of excess of chloride of barium does not affect its alkaline reaction.

Such of the sulphides as are soluble only in nitro-hydrochloric, or in concentrated nitric acid, become by the action of the acid converted into sulphates. That the sulphuric acid did not, however, exist in the original substance, is shown by fusing this latter with carbonate of sodium, when the fused mass will afford the reactions of a soluble sulphide instead of those of a soluble sulphate. Some sulphides are extremely difficult to dissolve completely in acid, in consequence of the deposition of sulphur, which fuses round the unaltered substance, and prevents any action of the acid upon it. This is particularly the case with the sulphides of arsenic and mercury in their ordinary sublimed state. But these sulphides are easily recognised by their colour, their volatility, and by their reducibility with conversion into sublimed metal, when heated with soda flux, for instance.

(61.) ACIDS OR SALTS OF HYDROGEN.

The ordinary solid acids are known by their solubility in cold or hot water to form strongly acid liquids, which effervesce with alkaline carbonates, are not precipitated by the several reagents for the metals, and do not evolve ammonia when heated with caustic potash. The boric and phosphoric acids yield, upon ignition, fusible residues of boric anhydride, and metaphosphoric acid respectively.

The oxalic, benzoic, tartaric, and citric acids are entirely dissipated by a prolonged heat, the two last furnishing an intermediate product of carbon.

The liquid acids, namely, the sulphuric, nitric, hydrochloric, and acetic acids, are known by their liquidity, volatility, strongly-marked acid reaction, and, except the sulphuric, by their characteristic odours. They give only negative results when tested for bases. The sulphuric and nitric acids are easily recognised by their respective actions on metallic copper, and hydrochloric acid by its action on peroxide of manganese (*vide* Chapter III.).

Acid salts, such, for instance, as the acid sulphates, oxalates, and tartrates of alkali-metal or ammonium, manifest a strongly

acid reaction to test-paper, and effervesce with alkaline carbonates. But they either evolve ammonia when treated with caustic potash, or leave a fixed residue upon ignition, which, in the case of the acid oxalates and tartrates, consists of alkaline carbonate. The acid salts of potassium react with perchloride of platinum, but not with tartaric acid, unless previously neutralised with soda,

(62.) MISCELLANEOUS SALTS.

Iodide of potassium solution reacts satisfactorily when treated with tartaric acid, but yields a dark brown liquid with perchloride of platinum. But after its iodine has been precipitated with excess of nitrate of silver, and the excess of silver with hydrochloric acid, the filtered liquid yields with perchloride of platinum the characteristic yellow precipitate of potassio-chloride of platinum.

Calomel occurs as a heavy, white, volatile powder. It is readily soluble in concentrated nitric acid, but in the act of solution becomes converted into a mercuric salt. That it was originally a mercurous salt is shown by the powder itself becoming black when agitated with potash-water.

White precipitate, amido-chloride of mercury, or chloride of mercurammonium, is a mercuric compound, readily distinguishable from calomel by heating it gently with potash-water, whereby it becomes of an orange-yellow colour, while ammoniacal vapours are given off. It is insoluble in water, soluble in nitric and hydrochloric acids, and dissipated by heat.

The sulphides of arsenic, known as realgar and orpiment, are orange or yellow coloured volatile solids. When boiled with nitro-hydrochloric acid, they are converted in great measure into arsenic acid, which may be obtained solid on evaporation. Its solution is precipitated very slowly by sulphuretted hydrogen, unless previously reduced by treatment with sulphurous acid.

(63.) LIQUID OR DISSOLVED SUBSTANCES.

A few drops of the liquid are evaporated on a glass slip, and the residue, if any, examined microscopically. Should there be an appreciable solid residue, a further quantity of liquid may be evaporated down in a capsule, the vapour examined for its odour and reaction, and the residue submitted to ignition, &c.; as described in pars. 25 and 26.

The reaction of the liquid to test-paper is next ascertained. Among volatile liquids, water, alcohol, and ether are neutral; ammonia, alkaline; and the sulphuric, nitric, hydrochloric, and acetic acids, strongly acid. These several liquids are readily distinguishable from one another by a few simple tests.

Solutions which leave a saline residue on evaporation, and are perfectly neutral, will most probably prove to contain some salt of an alkali- or alkaline earth- metal; salts of the heavy metals, with a few exceptions only, exhibiting a more or less marked acidity.

Solutions which have an alkaline reaction, known by their turning rose paper green, turmeric paper brown, or reddened litmus paper blue, may contain the hydrate or sulphydrate of an alkaline earth-metal; or the hydrate, sulphydrate, carbonate, phosphate, borate, or silicate of an alkali metal, or some heavy metal dissolved in excess of alkaline hydrate or carbonate (*vide* par. 64).

Solutions which have an acid reaction, known by their reddening blue litmus paper, may contain a free acid, an acid salt, or the normal salt of a heavy metal, in which last case the addition of even a drop of potash will most probably cause a precipitate.

After the above preliminary examination, the solution may be tested according to the directions of Tables I., II., III., and V.; or, in some cases, the evaporated residue may be dissolved in water or acid, and the solution so formed be employed by preference. The original solvent, if suspected to be other than water, may be distilled off, condensed in a receiver, and separately examined.

(64.) ALKALINE SOLUTIONS OF HEAVY METALS.

The hydrates of barium, strontium, and calcium, like those of the alkali metals, are soluble in water. The hydrates of all other metals are insoluble, and consequently, precipitable by caustic alkalies. The hydrates of lead, chromium, aluminum, and zinc, are readily soluble in excess of potash or soda; those of silver, copper, cadmium, nickel, cobalt, manganese, zinc, and magnesium readily, and those of chromium and iron (ferrous salts) sparingly soluble in excess of ammonia, especially in presence of neutral ammoniacal salts; while those of bismuth, mercury, and iron (ferric salts), are insoluble in excess of either reagent. On boiling alkaline solutions of chromium, the whole of the chromic hydrate is reprecipitated.

The hydrates of the metals tin, antimony, and arsenic, whose sulphides are soluble in sulphide of ammonium, have no practical interest in analysis. The tetroxide of arsenic is slightly soluble in water, and freely soluble in all alkaline liquids. The hydrate of antimony and both hydrates of tin are soluble in excess of potash, while stannic hydrate is soluble also in excess of ammonia. The potash solution of stannous hydrate is decomposed on boiling into metallic tin, which is deposited as a black powder, and stannate of potassium which remains dissolved.

The precipitates produced by carbonate of ammonium in solutions of silver, copper, nickel, cobalt, zinc, and magnesium, are readily soluble in excess of the reagent, especially when chloride of ammonium is also present, while those of iron (ferrous salts) and chrome are sparingly soluble. The precipitates produced in solutions of cadmium and manganese are insoluble in excess, as are also all those produced in solutions of metals whose hydrates are insoluble in ammonia. The precipitates produced by fixed alkaline carbonates are all insoluble in excess, except that produced in stannic salts, which dissolves in excess of the precipitant, and is again thrown down on ebullition.

§ VIII.—INDIVIDUAL BASES OF GROUP I.

The reactions of the individual bases of this group may be conveniently realised by operating with the following substances :

- TIN** . . The crystallised proto-chloride and precipitated peroxide.
ARSENIC . The white oxide.
ANTIMONY. The native tersulphide and tartar-emetic.
BISMUTH . The crystallised nitrate and precipitated oxide.
MERCURY . Corrosive sublimate, the red mercuric oxide, and the crystallised mercurous nitrate.
LEAD . . The oxide, carbonate, nitrate, and acetate.
SILVER. . The nitrate and oxide.
COPPER . The sulphate and oxide.
CADMIUM . The sulphate and carbonate.

(65.) TIN.

Tin salts are of two kinds, stannous or protosalts, represented by protochloride of tin SnCl_2 , and stannic or persalts, represented by perchloride of tin SnCl_4 .

a. When compounds of tin are heated upon charcoal with a mixture of carbonate of sodium and cyanide of potassium, a globule of white malleable metal is produced with very slight, if any, incrustation. If this globule be hammered out, and dissolved in hydrochloric acid, the tests for stannous salts can be applied to the solution so formed.

STANNOUS SALTS.

a. *Sulphydric acid* produces a brown precipitate of proto-sulphide of tin SnS , which dissolves in yellow sulphide of ammonium with conversion into persulphide of tin SnS_2 , so that on adding an acid to the solution, a yellow and not a brown precipitate is separated. The protosulphide of tin first thrown down is converted by boiling nitric acid into a white insoluble powder consisting of stannic anhydride SnO_2 .

β . Solution of *corrosive sublimate*, added carefully to stannous solutions, produces a white precipitate of calomel HgCl , which speedily becomes grey, and finally black, from its reduction to the state of metallic mercury.

STANNIC SALTS.

α . *Sulphydric acid* produces a yellow precipitate of disulphide of tin SnS_2 , which is insoluble in the carbonate, but soluble in the hydrate and sulphide of ammonium, and reprecipitable therefrom on the addition of an acid. It is likewise soluble in boiling hydrochloric acid, more readily on the addition of a little nitric acid also, and is converted by concentrated nitric acid into a white insoluble powder of stannic anhydride SnO_2 .

(66.) ARSENIC.

Arsenious acid is convertible into arsenic acid by boiling it with concentrated nitric acid, to which a little hydrochloric acid may be added with advantage. But in order to apply the various tests successfully, the acid liquid must be evaporated to dryness, and the residue dissolved in water. Arsenic acid is readily convertible into arsenious acid by the passage of sulphurous acid gas through its solution, or by heating it with sulphite of sodium and dilute hydrochloric acid.

α . *Sulphydric acid* produces, in acidulated solutions of arsenious acid, or of arsenic acid after the addition of sulphurous acid, a yellow precipitate of trisulphide of arsenic As_2S_3 , which is soluble in carbonate, hydrate, and sulphide of ammonium, and reprecipitated on the addition of an acid. It is insoluble in boiling hydrochloric acid, but is readily dissolved by hot nitric or nitro-hydrochloric acid.

β . *Nitrate of silver* produces in neutral or slightly ammoniacal solutions of arsenious acid a yellow precipitate of arsenite of silver Ag_3AsO_3 , and from similar solutions of arsenic acid a brick-dust red precipitate of arseniate of silver Ag_3AsO_4 . Both precipitates are soluble in excess of either ammonia or nitric acid.

γ. Sulphate of copper produces, in neutral or very faintly ammoniacal solutions of arsenious acid a grass-green precipitate of arsenite of copper $\text{HCu}''\text{AsO}_3$, and in similar solutions of arsenic acid a pale blue precipitate of arseniate of copper $\text{Cu}''_3(\text{AsO}_4)_4$. Both precipitates are soluble in excess of either ammonia or nitric or hydrochloric acid.

δ. When a compound of arsenic is mixed with soda-flux, and heated in a subliming tube, a steel-grey ring of reduced metal condenses in the upper or cool part of the tube.

(67.) ANTIMONY.

a. Sulphydric acid produces an orange-coloured precipitate of trisulphide of antimony Sb_2S_3 , which is insoluble in carbonate, but soluble in hydrate and sulphide of ammonium, and reprecipitable on the addition of an acid. It is also dissolved by hydrochloric acid with the aid of heat, and is converted almost entirely by strong nitric acid into a white insoluble powder, consisting of tetroxide of antimony Sb_2O_4 .

β. Water, added to certain antimony solutions (not to all), produces a white precipitate of a basic salt of antimony, soluble in excess of tartaric, hydrochloric, or nitric acid.

γ. Antimony compounds, when fused with carbonate of sodium on charcoal in the reducing flame, yield a bead of brittle metal with an abundant bluish-white incrustation. If the heat be prolonged, the metal volatilises entirely with the production of white fumes of tetroxide of antimony Sb_2O_4 .

(68.) BISMUTH.

a. Sulphydric acid produces a brownish-black precipitate of trisulphide of bismuth Bi_2S_3 , which is insoluble in sulphide of ammonium. It dissolves readily in hot nitro-muriatic, nitric, or hydrochloric acid.

β. Caustic alkalies give a white precipitate of hydrate of

bismuth $\text{Bi}_2\text{O}_3 \cdot \text{H}_2\text{O}$, insoluble in excess of either potash or ammonia.

γ . *Water*, when added to moderately-concentrated and not over-acid solutions of bismuth, causes a dense white precipitate of a basic bismuth salt, which does not disappear on the addition of tartaric acid, but dissolves in excess of nitric or hydrochloric acid.

δ . Bismuth compounds, when mixed with carbonate of sodium and heated upon charcoal in the reducing blowpipe flame, yield a brittle metallic globule and a yellow incrustation.

(69.) MERCURY.

There are two classes of mercury salts, namely, the mercurous, represented by calomel Hg_2Cl_2 , and the mercuric, represented by corrosive sublimate HgCl_2 . Certain reactions are common to both, while others are distinctive between them.

α . An excess of *sulphydric acid* produces a black precipitate of mercurous sulphide Hg_2S , or of mercuric sulphide HgS , which is insoluble in sulphide of ammonium, and also in strong hot nitric and hydrochloric acids taken separately, but is readily soluble in a mixture of the two. An insufficiency of sulphydric acid produces in mercuric salts a highly characteristic white precipitate of doubtful composition, which, as the proportion of gas increases, becomes orange, brown, and finally black.

β . *Protochloride of tin* produces at first a white precipitate of calomel Hg_2Cl_2 , which becomes in succession grey and almost black on adding more of the reagent and warming gently. If the supernatant liquor be poured off, and the deposit boiled with hydrochloric acid, globules of metallic mercury make their appearance.

γ . Mercury compounds mixed with carbonate of sodium, and heated in a reduction-tube, furnish a sublimate of well-defined mercurial globules.

MERCUROUS SALTS.

a. Potash and *ammonia* alike produce a black precipitate, the former of mercurous hydrate $\text{Hg}_2\text{O} \cdot \text{H}_2\text{O}$, the latter of some amido-mercurous compound.

β. Hydrochloric acid produces a white precipitate of calomel Hg_2Cl_2 , which dissolves in boiling nitric acid. It is turned black by ammonia, being converted into amido-mercurous chloride.

MERCURIC SALTS.

a. Potash gives a yellow precipitate of mercuric oxide HgO , which is turned white on the addition of ammonia, or in presence of an ammoniacal salt. *Ammonia* produces a white precipitate of some amido-mercuric compound.

β. Iodide of potassium, added carefully, produces a bright orange-red precipitate of mercuric iodide HgI_2 , which dissolves in excess of the reagent, forming a colourless solution.

(70.) LEAD.

a. Sulphydric acid produces a black (occasionally red) precipitate of sulphide of lead PbS , which is insoluble in sulphide of ammonium, but dissolves when heated with not too concentrated nitric or hydrochloric acid. Strong nitric acid converts it into a white insoluble deposit of sulphate of lead PbSO_4 .

β. Caustic alkalies give a white precipitate of hydrate of lead $\text{PbO} \cdot \text{H}_2\text{O}$, soluble in excess of potash.

γ. In moderately strong solutions, hydrochloric acid gives a white crystalline precipitate of chloride of lead PbCl_2 , which is soluble in boiling water, unaffected by ammonia, and soluble in great excess of potash.

δ. Dilute sulphuric acid or a dissolved sulphate gives a dense white precipitate of sulphate of lead PbSO_4 , which is insoluble in dilute acids, soluble in strong hydrochloric acid with the aid of heat, and also in a large excess of potash.

ε. Lead compounds, when fused with carbonate of sodium and charcoal in the reducing blowpipe flame, yield a globule of soft metal and a brownish-yellow incrustation.

(71.) SILVER.

α. *Sulphydric acid* gives a black precipitate of sulphide of silver Ag_2S , insoluble in sulphide of ammonium, soluble in warm nitric acid, and converted by ebullition with hydrochloric acid into a white deposit of chloride of silver AgCl .

β. *Caustic alkalies* give a brown precipitate of hydrate of silver $\text{Ag}_2\text{O} \cdot \text{H}_2\text{O}$, which is insoluble in excess of potash, but soluble in ammonia, forming a colourless solution.

γ. *Hydrochloric acid* gives in solutions of silver salts a white precipitate of chloride of silver AgCl , which is insoluble even in boiling nitric acid, but readily soluble in ammonia. The colour of the precipitate changes to a slate-purple by exposure to light.

δ. Silver compounds, when fused with carbonate of sodium upon a charcoal support in the reducing blowpipe flame, yield a button of hard white malleable metal, without any incrustation being formed on the charcoal.

(72.) COPPER.

α. *Sulphydric acid* produces a dark-brown precipitate of sulphide of copper CuS , which is insoluble in sulphide of potassium, and but sparingly soluble in sulphide of ammonium. It dissolves readily in nitric but not in hydrochloric acid, save by an application of heat.

β. *Potash* gives a pale blue precipitate of hydrate of copper $\text{CuO} \cdot \text{H}_2\text{O}$, which is insoluble in excess, and converted by ebullition into black oxide of copper CuO . *Ammonia* gives a similar blue precipitate of hydrate of copper, which is soluble in excess of the reagent, forming a deep blue solution, the transparency of which is not affected by the addition of a moderate quantity of potash.

The pale-blue precipitate produced by *carbonate of ammonium* is also readily soluble in excess of the reagent, with production of a deep blue liquid.

γ. *Ferrocyanide of potassium* gives a chocolate-coloured precipitate of ferrocyanide of copper Cu_2FeCy_6 , which is decomposed by caustic potash, and is then freely soluble in ammonia, forming a deep blue liquid.

δ. A piece of clean *iron* or steel dipped into an acidulated copper solution becomes coated with metallic copper, which may be dissolved off by the conjoint action of ammonia and air into a deep blue liquid.

ε. The borax bead heated with a particle of any copper-compound becomes blue or green in the oxidising, and nearly colourless or reddish-grey in the reducing, flame.

(73.) CADMIUM.

α. The precipitate of sulphide of cadmium CdS , produced by *sulphydric acid*, is of a bright yellow colour, and insoluble in sulphide of ammonium. It disappears readily on the addition of nitric or hydrochloric acid, and does not form in very acid solutions.

β. *Caustic alkalies* give a white precipitate of hydrate of cadmium $\text{CdO} \cdot \text{H}_2\text{O}$, soluble in excess of ammonia, but not in that of potash. The precipitate produced by carbonate of ammonium does not disappear in excess of the reagent.

γ. Cadmium compounds, fused with carbonate of sodium in the reducing blowpipe flame, give a reddish-brown incrustation of oxide of cadmium CdO , but no bead of metal.

§ IX.—INDIVIDUAL BASES OF GROUP II.

The reactions of the individual bases of this group may be conveniently realised by operating with the following substances:

NICKEL . . . The oxide and sulphate.

COBALT . . . The oxide.

MANGANESE. The sulphate, chloride, and carbonate.

IRON . . . The red and black oxides, the sulphate and sulphide.

ZINC . . . The oxide and sulphate.

CHROMIUM . The precipitated oxide and chrome alum.

ALUMINIUM . The hydrate and sulphate.

(74.) NICKEL.

Solutions of nickel are generally of a green colour.

a. Sulphide of ammonium gives with nickel salts a black precipitate of sulphide of nickel NiS , not soluble in hydrochloric acid until after the addition of a drop or two of nitric acid.

β. Ammonia gives a slight greenish precipitate of hydrate of nickel $\text{NiO.H}_2\text{O}$, soluble in excess of the reagent, forming a violet-blue liquid, from which potash reprecipitates the green hydrate. Moreover, *potash* throws down this hydrate readily from all ordinary nickel solutions.

γ. In the reducing flame, nickel renders the borax bead purplish-grey and turbid, and in the oxidising flame of a dark sherry colour with a tinge of violet. If a fragment of nitre be added, and the bead again heated in the oxidising flame, a well-marked purple colour is produced.

(75.) COBALT.

Cobalt solutions are pink when dilute, blue when concentrated.

a. Sulphide of ammonium gives with cobalt salts a black precipitate of sulphide of cobalt CoS , not soluble in hydrochloric acid until after the addition of a few drops of nitric acid.

β. Potash gives a blue precipitate of prothydrate of cobalt $\text{CO}_2\text{H}_2\text{O}$, insoluble in excess of the precipitant. Excess of *ammonia*, especially in presence of chloride of ammonium, produces a brownish-pink solution, becoming brown and opaque on exposure to air, from the deposition of the insoluble sesquihydrate.

γ. Cobalt compounds impart to the borax bead, when heated in either flame of the blowpipe, a deep sapphire-blue colour.

(76.) MANGANESE.

Manganese solutions are of a faint pink tinge, or altogether colourless.

a. Sulphide of ammonium gives a buff-coloured precipitate of sulphide of manganese MnS , soluble even in acetic acid.

β. Potash gives a white precipitate of prothydrate of manganese $\text{MnO}_2\text{H}_2\text{O}$, speedily becoming brown on exposure, from its conversion into the sesquihydrate $\text{Mn}_2\text{O}_3\text{H}_2\text{O}$. If to the solution of a manganous salt containing chloride of ammonium, an excess of *ammonia* be added at once, a clear colourless solution will be produced, quickly becoming brown and opaque when exposed to the air, from a conversion of the soluble proto- into the insoluble sesqui-compound.

γ. The borax bead with manganese is amethyst-red in the oxidising, and nearly colourless in the reducing, flame. Its appearance is much interfered with by the presence of a trace of iron.

δ. Compounds of manganese, fused upon platinum foil with *carbonate of sodium* in the oxidising flame of the blowpipe, either with or without the addition of a little nitre, yield a bluish-green fusible mass of manganate of sodium Na_2MnO_4 . For this experiment but a very small quantity of the manganese compound should be employed.

(77.) IRON.

There are two classes of iron salts, namely, ferrous or proto-salts, represented by green vitriol FeSO_4 , and ferric or persalts, represented by sesquichloride of iron FeCl_3 or Fe_2Cl_6 . The proto-salts are rarely ever free from some admixture with persalts.

a. Sulphide of ammonium gives a black precipitate of protosulphide of iron FeS , readily soluble in hydrochloric acid. The precipitate, when thrown down from ferric salts, is accompanied by free sulphur.

β. Iron compounds dissolve in the borax bead, forming a glass which is of a bottle-green colour in the reducing, and of a yellowish-brown colour in the oxidising, flame. Ferrous compounds give the green, and ferric compounds the yellow, colour most readily.

FERROUS SALTS.

Ferrous solutions are generally colourless, or of a pale green tint.

a. Caustic alkalis give a dingy green precipitate of ferrous hydrate $\text{FeO} \cdot \text{H}_2\text{O}$, which becomes red on exposure to air. The pure, nearly white prothydrate is sparingly soluble in excess of ammonia and chloride of ammonium.

β. Ferrocyanide of potassium gives a pale blue precipitate of double ferrocyanide of iron and potassium $\text{K}_4\text{Fe}_2\text{Cy}_6$, which becomes dark blue on exposure to air.

γ. Ferridcyanide of potassium gives a dark-blue precipitate of ferridcyanide of iron $\text{Fe}_5\text{Cy}_{12}$, or $\text{Fe}'''_3\text{Fe}_2\text{Cy}_{12}$, insoluble in dilute acids and decomposable by caustic alkalis.

FERRIC SALTS.

Ferric solutions are generally of a yellow, brown, or red colour.

a. Caustic and carbonated alkalis alike give a brick-dust red precipitate of ferric hydrate $\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$, accompanied in the case

of the carbonates by an evolution of carbonic acid. The precipitate is quite insoluble in the hydrates of potassium and ammonium, and also in their respective carbonates at a boiling heat.

β . *Ferrocyanide of potassium* gives a deep blue-coloured precipitate of sesquiferrocyanide of iron $\text{Fe}_7\text{Cy}_{18}$, or $\text{Fe}^{\text{III}}_3\text{Fe}^{\text{II}}_4\text{Cy}_{18}$, insoluble in dilute acids, and decomposable by caustic alkalies. The ferridcyanide does not produce any precipitate, but gives to the ferric solution a colour which is brown or red according to circumstances.

γ . *Sulphocyanate of potassium* imparts to ferric solutions a deep red colour, from the formation of ferric sulphocyanate FeCy_3S_3 , or $\text{Fe}(\text{CyS})_3$.

(78.) ZINC.

Zinc solutions are usually colourless.

α . *Sulphide of ammonium* gives a white precipitate of sulphide of zinc ZnS , which is insoluble in acetic, but readily soluble in hydrochloric acid. Unlike most zinc precipitates, it is insoluble in any alkaline solution.

β . *Caustic alkalies* give with solutions of zinc a white precipitate of hydrate of zinc $\text{ZnO} \cdot \text{H}_2\text{O}$, soluble in a large excess of the precipitant. *Alkaline carbonates* also give a white precipitate of a highly basic carbonate of zinc, insoluble in excess of the fixed alkaline carbonates, but soluble in presence of ammoniacal salts.

γ . Zinc salts, heated on platinum foil, leave a fixed infusible residue, which is yellow when hot, white when cold. After being moistened with solution of nitrate of cobalt, and reignited in the blowpipe flame, it assumes a fine green colour.

δ . Zinc salts, mixed with carbonate of sodium or cyanide flux, and heated in the reducing blowpipe flame, deposit upon the charcoal an incrustation which is yellow when hot, white when cold, and green after ignition with nitrate of cobalt.

(79.) CHROMIUM.

Solutions of chromic salts are mostly of a green or violet colour.

a. Caustic alkalies, their *sulphides* or *sulphydrates*, and *carbonates* throw down from chrome solutions a greenish precipitate of hydrate of chrome $\text{Cr}_2\text{O}_3 \cdot \text{H}_2\text{O}$, which is slightly soluble in excess of ammonia, more so in excess of potash, and in each instance reprecipitable on boiling. Its production by sulphides and carbonates respectively is accompanied by an evolution of sulphydric or carbonic acid.

β. Chrome imparts to the borax bead a fine green colour, permanent in both flames of the blowpipe.

γ. When a chrome compound is fused on platinum foil with a little carbonate of sodium and nitre, a yellow mass of chromate of sodium Na_2CrO_4 , is produced, which dissolves in water to form a solution, giving with acetate of lead a yellow precipitate of chromate of lead PbCrO_4 .

(80.) ALUMINUM.

a. Caustic alkalies, their *carbonates* and *sulphides* or *sulphydrates*, give with aluminous solutions a white gelatinous precipitate of hydrate of aluminum $\text{Al}_2\text{O}_3 \cdot \text{H}_2\text{O}$, which is soluble in excess of potash, but reprecipitated on the addition of chloride of ammonium. The action of the carbonates and sulphides respectively is accompanied by an evolution of carbonic or sulphydric acid.

β. Salts of aluminum, when heated on charcoal or platinum foil, leave a white, infusible, highly incandescent residue, which, when moistened with nitrate of cobalt solution and reignited in the blowpipe flame, assumes a bright blue colour.

§ X.—INDIVIDUAL BASES OF GROUP III.

The reactions of the individual bases of this group may be conveniently realised by operating with the following substances:

BARIUM	. . The carbonate, nitrate, and chloride.
STRONTIUM	. The carbonate and nitrate.
CALCIUM	. The hydrate, carbonate, phosphate, oxalate, and sulphate.
MAGNESIUM	. The oxide, carbonate, and sulphate.
POTASSIUM	. The nitrate and sulphate.
SODIUM	. . The carbonate, sulphate, chloride, and phosphate.
AMMONIUM	. The sulphate, chloride, carbonate, phosphate, and oxalate.

(81.) BARIUM, STRONTIUM, CALCIUM.

These three metals possess many properties in common.

α. Their hydrates $MO.H_2O$ or $M(HO)_2$, are all soluble in water, so that the addition of caustic alkali, if perfectly free from carbonate, does not disturb their solutions. Their sulphides and sulphhydrates are also soluble.

β. The following salts of these metals are insoluble in water, namely, the oxalates, phosphates, carbonates (even in the presence of ammoniacal salts), and sulphates, sulphate of barium being the most insoluble, sulphate of calcium the least so. Hence neutral barium, strontium, and calcium salts are precipitated by soluble oxalates, phosphates, carbonates and sulphates.

γ. When a barium, strontium, or calcium salt is ignited on platinum foil, a white fixed residue remains, which, except in the case of the chlorides, is usually infusible. When moistened with solution of nitrate of cobalt and reignited, the infusible mass acquires an ill-defined grey colour.

(82.) BARIUM.

a. *Sulphuric acid* and solutions of all *sulphates*, even when very dilute, give with barium salts a white precipitate of sulphate of barium $BaSO_4$, insoluble in acids and alkalies.

β. Hydrofluosilicic acid produces in acid and neutral solutions a somewhat transparent precipitate of fluosilicate of barium BaSiF_6 , the deposition of which is much facilitated by the addition of a little alcohol.

γ. Barium salts, when intensely heated before the blowpipe upon a fine platinum wire, impart a marked apple-green colour to the flame.

(83.) STRONTIUM.

α. Sulphuric acid and solutions of most *sulphates*, give a precipitate of sulphate of strontium SrSO_4 , insoluble in acids and alkalis, very sparingly soluble in water. Strontium salts are not precipitated by solution of sulphate of strontium, and very slowly by solutions of the sulphates of calcium and potassium.

β. Strontium salts, when intensely heated before the blowpipe upon a fine platinum wire, impart a deep crimson colour to the flame.

(84.) CALCIUM.

α. Sulphuric acid and strong solutions of *sulphates* give with moderately strong calcium solutions a white precipitate of sulphate of calcium CaSO_4 , slightly soluble in water, insoluble in proof-spirit. But dilute calcium solutions are not precipitable by any sulphate; while even strong solutions are not precipitable by the sulphates of strontium, calcium, and scarcely by that of potassium.

β. Oxalate of ammonium produces in neutral calcium solutions a white precipitate of oxalate of calcium CaC_2O_4 , soluble in nitric and hydrochloric acids, insoluble in acetic and oxalic acids, and insoluble in ammonia.

γ. Calcium salts impart a fine orange-red colour to the blow-pipe flame.

(85.) MAGNESIUM.

α. Sulphhydrate and sulphate of magnesium are soluble in water. Hence magnesian salts are not precipitated by soluble sulphhydrates and sulphates.

β . Hydrate, carbonate, oxalate, phosphate, and arseniate of magnesium, are insoluble in water. Hence magnesian salts are precipitated by soluble hydrates, carbonates, oxalates, phosphates, and arseniates. But all magnesian precipitates, excepting the phosphate and arseniate, are soluble in ammoniacal solutions. When *phosphate of sodium* or *ammonium* is added to a solution of magnesium containing chloride of ammonium and rendered alkaline by ammonia, a white crystalline precipitate of triple phosphate $\text{Mg}(\text{NH}_4)\text{PO}_4$ is produced, which is soluble in dilute acids, insoluble in ammonia and ammoniacal salts.

γ . Magnesium compounds, heated on charcoal or platinum foil, leave a white fixed infusible residue, which when moistened with nitrate of cobalt solution and reignited in the blowpipe flame assumes a faint pink colour.

(86.) POTASSIUM.

α . All simple potassium salts, except the acid-tartrate, are moderately soluble in water. *Tartaric acid* added in excess to the neutral or alkaline solution of a potassium salt throws down a white crystalline precipitate of cream of tartar $\text{KH}_5\text{C}_4\text{O}_6$, which frequently does not appear immediately. Its deposition is facilitated by stirring the mixed liquids, and by the addition of a little alcohol to them. It is soluble in mineral acids, in hot water, and in a large excess of cold water.

β . *Perchloride of platinum* when added to neutral or acid solutions of potassium throws down a yellow crystalline precipitate of potassio-chloride of platinum K_2PtCl_6 or $2\text{KCl}.\text{PtCl}_4$. The solution should generally be acidulated with a few drops of hydrochloric acid before being tested. The precipitate does not always appear immediately: its production is facilitated by stirring and by the addition of a little alcohol.

γ . Potassium salts when heated on charcoal or platinum foil leave a fixed and generally fusible residue. When heated before the blowpipe they impart to the flame a marked violet colour,

which, however, is not recognisable in the presence of even a small quantity of sodium.

(87.) AMMONIUM.

α . All simple ammoniacal salts, except the acid tartrate, are freely soluble in water. *Tartaric acid* reacts upon ammoniacal as upon potassium salts. The resulting precipitate of acid tartrate of ammonium $(\text{NH}_4)_2\text{H}_2\text{C}_4\text{O}_6$, is, however, more soluble in water than the corresponding potassium compound, and consequently does not form, save when the ammoniacal solution is moderately concentrated.

β . *Perchloride of platinum* reacts upon ammoniacal salts to form the ammonio-chloride of platinum $(\text{NH}_4)_2\text{PtCl}_6$, or $2\text{NH}_4\text{Cl}.\text{PtCl}_4$, which closely resembles the corresponding potassium compound in its appearance, solubility, and mode of deposition.

γ . When any ammoniacal salt is boiled with *potash* or *lime*, ammoniacal vapour is given off, which is recognisable by its smell, by its action on test paper, and by its forming opaque fumes when brought into contact with the vapour of hydrochloric acid.

δ . All ammoniacal salts volatilise partly, most of them entirely, when heated upon platinum foil or charcoal.

(88.) SODIUM.

α . Sodium solutions are non-precipitable, all simple sodium salts being soluble in water.

β . Sodium salts when heated on platinum foil or charcoal leave a fixed residue, almost always fusible, and impart an intense yellow colour to flame.

§ XL.—REACTIONS OF INDIVIDUAL ACIDS.

(89.) CHROMATES.

Chromates are usually of a yellow or red colour, and, except those of the alkali metals, are more or less insoluble in water.

a. Chromates acidified with hydrochloric acid, and treated with *sulphuretted hydrogen*, yield a deposit of sulphur, and a green solution of chromic chloride, CrCl_3 or Cr_2Cl_6 . The same compound is produced on boiling the solution of a chromate with hydrochloric acid and a little alcohol. To it the ordinary tests for chrome salts can be applied.

β. Nitrate of silver gives a dark red precipitate of chromate of silver Ag_2CrO_4 , soluble in nitric acid. Nitrate of barium and acetate of lead throw down yellow precipitates of the chromates of the respective metals.

(90.) NITRATES.

The nitrates are soluble or non-precipitable salts, which de-flagrate when ignited upon charcoal or with organic matter.

a. When a nitrate, or the concentrated solution of a nitrate, is gently heated with *sulphuric acid* and a few *copper turnings*, brown fumes of peroxide of nitrogen N_2O_4 , are evolved, which redden, but do not bleach, litmus paper, and produce a purple colouration on starch paper moistened with iodide of potassium.

Fig. 36.

β. A solution of *protosulphate of iron* carefully poured upon *sulphuric acid*, to which a minute portion of a solid or dissolved nitrate has been added, produces a deep brown halo at the junction of the two liquids, as seen in fig. 36.

γ. When a minute quantity of *gold-leaf* is boiled in *hydrochloric acid*, no action is produced; but, on the addition of a little nitric acid or a nitrate, the gold quickly disappears, and may be detected in solution by protochloride of tin, which gives rise to a purplish precipitate.



(91.) CHLORATES.

The chlorates, like the nitrates, are soluble or non-precipitable

salts, which deflagrate when ignited upon charcoal or with organic matter.

a. Strong *sulphuric acid* added to a solid chlorate produces a brown colouration with crackling detonation, especially when gently warmed. The experiment should be made carefully, and with but a small quantity of the salt, for fear of an explosion and scattering of the acid.

β. The addition of some solid or dissolved chlorate to *hydrochloric acid* enables it to bleach litmus and dissolve gold-leaf. Strong hydrochloric acid reacts with a moderate quantity of a solid chlorate to produce a greenish-yellow gas known as *euchlorine*.

γ. Chlorates of alkali-metal, when heated alone, and other chlorates, when heated with carbonate of sodium in a reduction tube, evolve oxygen, known by its inflaming a piece of incandescent wood; and leave a residue of alkaline chloride, recognisable by adding nitrate of silver to its solution previously acidified with nitric acid, when a white precipitate of chloride of silver is thrown down.

(92.) CARBONATES.

On adding *hydrochloric* or slightly diluted *sulphuric acid* to a solid or dissolved carbonate, effervescence of carbonic anhydride CO_2 , is produced, either immediately or on gently warming. The gas is free from any marked smell, and does not affect lead paper, but turns *lime water* milky, from the precipitation of carbonate of calcium CaCO_3 . It may be tested by means of the apparatus shown in fig. 37; or, being one and a half times as heavy as air, it may be generated in one test tube, and then carefully poured into another containing some lime water, with which it is to be agitated. The lime water may be replaced by *baryta-water* or a solution of basic acetate of lead.

β. Soluble carbonates give precipitates with salts of silver, barium, calcium, &c., &c., which disappear with effervescence on the addition of any acid.

(93.) SULPHIDES AND SULPHYDRATES.

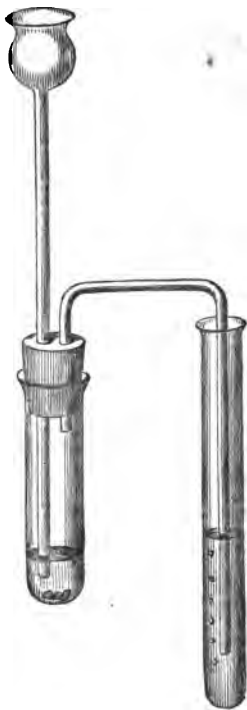
α. Most sulphides when acted upon by hydrochloric or moderately dilute sulphuric acid, especially on the application of heat, evolve sulphuretted hydrogen, which is known by its offensive smell, and by the brown or lustrous black stain it produces on lead paper. All sulphides, when fused with carbonate of sodium in a porcelain capsule, leave a residue which disengages sulphuretted hydrogen upon the addition of an acid. The residue, moistened with water, gives a black stain to silver coin, and a purple colouration with nitroprusside of sodium (vide β and γ).

β. Soluble sulphides, including sulphide of hydrogen, give with *nitrate of silver* a black precipitate of sulphide of silver Ag_2S , which does not disappear on the addition of cold nitric acid. They also produce a black stain on metallic silver, or *acetate of lead* paper. The precipitate of sulphide of lead PbS , thrown down from any dissolved lead salt, is not affected by dilute mineral acids, save on the application of heat.

γ. *Nitroprusside of sodium* produces a magnificent purple colouration with soluble sulphides, even when extremely dilute. The effect, however, is not obtained with sulphide of hydrogen until after the addition of a drop or so of alkali.

δ. Many sulphides, when heated in a glass tube open at both ends, evolve sulphurous anhydride SO_2 , which is known by its

Fig. 37.



smell of burning sulphur, and by its producing a purple stain on starch paper moistened with iodic acid.

(94.) SULPHITES.

a. Hydrochloric or sulphuric acid causes an effervescence of sulphurous acid or anhydride SO_2 , known by its smell of burning sulphur, and by the purple colouration it gives to starch paper moistened with iodic acid. On the addition of a fragment of zinc, the sulphurous is replaced by sulphydric acid, known by its effect on lead paper. Hyposulphites behave in a similar manner, except that the effervescence of sulphurous acid is accompanied by a deposition of yellow sulphur.

β . Perchloride of iron produces in neutral solutions a red colouration, which disappears on the addition of a strong acid, and is destroyed by boiling, with formation of a brown deposit of ferric oxide.

(95.) SULPHATES.

a. Nitrate or chloride of barium throws down an opaque white precipitate of sulphate of barium BaSO_4 , which does not disappear on the addition of hydrochloric or nitric acid even at a boiling heat. But in presence of a large excess of nitric acid a portion of sulphate of barium remains in solution. The seleniate and silicofluoride of barium resemble the sulphate in being precipitable from acid solutions; but, when boiled with strong hydrochloric acid, the seleniate dissolves with evolution of chlorine, while the silicofluoride, which is very transparent, dissolves, with the exception of a little silica.

(96.) CHLORIDES.

a. Nitrate of silver produces a white clotty precipitate of chloride of silver AgCl , which becomes slate-coloured on exposure to light. It is insoluble even in boiling concentrated nitric acid, but is readily soluble in ammonia. When heated in a

porcelain capsule, it does not undergo decomposition, but simply fuses.

β . Chlorides, when heated with strong *sulphuric acid*, save those of mercury, silver, and tin, evolve hydrochloric acid gas, known by its pungent smell and acid reaction. When heated with *peroxide of manganese* and sulphuric acid, chlorine gas is given off, recognisable by its irritant odour, green colour, power of bleaching litmus, &c., and by the purple stain it produces on a piece of starch paper moistened with iodide of potassium.

(97.) BROMIDES.

α . The precipitate of bromide of silver AgBr , produced by *nitrate of silver* in solutions of bromides, closely resembles the chloride, except that it has a faint tinge of yellow and is less readily soluble in ammonia.

β . Bromides when heated with strong *sulphuric acid*, with or without the addition of *peroxide of manganese*, evolve bromine, which is recognised by the red colour and irritant smell of its vapour, by its bleaching litmus, and turning starch paper yellow.

(98.) IODIDES.

α . The precipitate of iodide of silver AgI , produced by *nitrate of silver* in solutions of iodides, is of a pale yellow colour, insoluble in, but turned white by, ammonia, and in other respects similar to the chloride.

β . A sufficiency of *chloride of mercury* produces a scarlet precipitate of iodide of mercury HgI_2 ; while *acetate of lead* throws down a yellow precipitate of iodide of lead PbI_2 , somewhat soluble in boiling water, and deposited therefrom on cooling, in golden scales.

γ . Iodides when acted upon by strong *sulphuric acid* with or without *peroxide of manganese*, evolve iodine, known by its violet vapour, staining starch paper purple.

δ. A drop or two of *chlorine water*, or of nitric or preferably nitro-hydrochloric acid, added to a dissolved iodide, causes a liberation of its iodine, which, if in any quantity, imparts a yellow-brown colour to the liquid. But in presence of dilute starch paste, a deep purple colour is developed; and chloroform or disulphide of carbon shaken with the liquid becomes of a bright pink or crimson colour.

(99.) CYANIDES.

α. *Nitrate of silver* throws down a white clotty precipitate of cyanide of silver AgCl , not altered in colour by exposure to light. It is readily soluble in ammonia, insoluble in cold, but soluble in boiling concentrated nitric acid.

The white precipitates produced by nitrate of silver in ferrocyanide and sulphocyanate solutions, and the brown precipitate produced in ferridcyanide solutions are insoluble in nitric acid.

β. When a mixture of ordinary *sulphate of iron* and *potash* is added to a simple cyanide, no obvious effect is produced, but on acidification with hydrochloric acid the unaltered hydrates of iron are dissolved, and Prussian blue or sesquiferrocyanide of iron, $\text{Fe}_7\text{Cy}_{18}$ or $\text{Fe}_3'''\text{Cy}_6$, $\text{Fe}_4'''\text{Cy}_{12}$, is left as a deep blue precipitate.

γ. Cyanides, including cyanide of silver, when acted upon by yellow *sulphide of ammonium*, are converted into sulphocyanates MCyS_2 . After evaporating to dryness, to expel the excess of sulphide of ammonium, *perchloride of iron* gives a deep red colouration due to the production of ferric sulphocyanate $\text{Fe}'''\text{(CyS)}_3$.

(100.) PHOSPHATES.

α. *Sulphate of magnesium* added to a phosphate solution rendered alkaline by ammonia, and containing chloride of ammonium, produces a white crystalline precipitate of phosphate of magnesium and ammonium $\text{Mg}(\text{NH}_4)\text{PO}_4$, readily soluble in acids. The formation of the precipitate is facilitated by rubbing the

inside of the tube with a stirring rod. *Chloride of barium* or *calcium* produces in neutral or ammoniacal phosphate solutions a white precipitate of phosphate of calcium or barium, soluble even in acetic acid.

β . *Nitrate of silver* added to a neutral or nearly neutral phosphate solution, produces a yellow (under certain circumstances white) precipitate of phosphate of silver Ag_3PO_4 , soluble in ammonia and dilute nitric acid.

γ . *Perchloride of iron* produces in phosphate solutions which are neutral, or acid only with acetic acid, a pale brown precipitate of phosphate of iron Fe^+PO_4 . An excess of acetate of ammonium may be added to a solution of phosphate of magnesium or calcium in the smallest sufficient quantity of hydrochloric acid, whereby the free hydrochloric is replaced by free acetic acid. Then, on adding a drop of perchloride of iron, a deep red colour, speedily changing into an opaque white cloud, is produced. On adding more perchloride of iron gradually, until the liquid remains permanently red, and afterwards boiling and filtering, a colourless filtrate is obtained, free alike from phosphoric acid and iron, but containing chloride of calcium or magnesium.

δ . Excess of *molybdate of ammonium* added to a phosphate solution containing free nitric acid, produces, either immediately or on gentle warming, a bright yellow precipitate of a phosphomolybdate of ammonium of uncertain composition.

Except that the arseniate of silver is brick red instead of yellow or white, all the above described reactions apply equally to arsenic and phosphoric acids. Arsenic acid may, however, be got rid of by treatment of its solution with sulphurous acid and sulphydric acid in succession, or by prolonged treatment with sulphydric acid alone (vide par. 66).

(101.) OXALATES.

Unlike most organic acids, save those which volatilise without decomposition, neither oxalic acid nor its salts become charred by

the action of heat or strong sulphuric acid. They effervesce with dilute sulphuric acid and peroxide of manganese.

a. Oxalates soluble in water, or oxalates soluble in acid after the addition of acetate of ammonium, give with *chloride* or even *sulphate of calcium* a white precipitate of oxalate of calcium CaC_2O_4 , insoluble in ammonia and in acetic acid, but soluble in dilute mineral acids.

β. *Nitrate of silver* and *chloride of barium* give with neutral oxalates white precipitates of oxalate of silver $\text{Ag}_2\text{C}_2\text{O}_4$, and oxalate of barium BaC_2O_4 , respectively, soluble in dilute nitric, but insoluble or very sparingly soluble in acetic acid.

(102.) TARTRATES.

Tartaric acid and tartrates, when ignited, evolve a peculiar odour, and leave an abundant charcoal. They become blackened when heated with strong sulphuric acid; whereas citric acid and the citrates acquire only a yellow colour.

a. *Chloride of calcium* produces in neutral tartrates a white precipitate of tartrate of calcium $\text{Ca}^+\text{H}_4\text{C}_4\text{O}_6^-$, soluble even in acetic acid. On heating the washed precipitate with aqueous potash, filtering the mixture and boiling the filtrate, a white turbidity is produced which disappears on cooling. Chloride of calcium does not precipitate neutral citrates in the cold; but, upon boiling, a temporary white turbidity is produced.

β. *Nitrate of silver* throws down from neutral tartrates a white precipitate of tartrate of silver $\text{Ag}_2\text{H}_4\text{C}_4\text{O}_6$, soluble in dilute nitric acid and in ammonia, and decomposed on boiling, with blackening and reduction. *Chloride of barium* does not precipitate tartrates in the presence of ammoniacal salts or free acid.

γ. Neutral tartrates of potassium, and of sodium and ammonium in presence of *chloride of potassium*, yield, on acidification with *acetic acid*, a white crystalline precipitate of cream of tartar, or acid tartrate of potassium $\text{KH}_3\text{C}_4\text{O}_6$ which deposits

most readily on stirring, and is soluble in boiling water and mineral acids.

(103.) ACETATES.

a. Perchloride of iron added to a neutral acetate produces a deep red colour, from the formation of peracetate of iron $\text{Fe}^{\text{III}}(\text{H}_3\text{C}_2\text{O}_4)_3$. The colour is destroyed on the addition of a mineral acid, or on boiling; in the latter case with formation of an ochry deposit.

β. Nitrate of silver produces, in concentrated solutions only, a white precipitate of acetate of silver $\text{AgH}_3\text{C}_2\text{O}_4$, soluble in hot water, in dilute nitric acid, and in ammonia.

γ. On warming a solid acetate, or its concentrated solution, with sulphuric acid, acetic acid is liberated, recognisable by its well-known smell; but in presence of *alcohol* the fragrant vapour of acetic ether $(\text{C}_2\text{H}_5)_2\text{H}_3\text{C}_2\text{O}_4$, is evolved instead.

(104.) BENZOATES.

Benzoates evolve the characteristic smell of benzene upon ignition, and yield a sublimate of benzoic acid when warmed with strong sulphuric acid; but in neither case do they undergo much charring.

a. Perchloride of iron gives with neutral benzoates a brownish-yellow precipitate of perbenzoate of iron $\text{Fe}^{\text{III}}(\text{H}_5\text{C}_7\text{O}_4)_3$, which dissolves completely in warm dilute hydrochloric acid forming a solution from which benzoic acid crystallises out on cooling.

β. Nitrate of silver gives a white clotty precipitate of benzoate of silver $\text{AgH}_5\text{C}_7\text{O}_4$, soluble in hot water, in dilute nitric acid, and in ammonia.

(105.) BORATES.

Borate solutions have usually an alkaline reaction to test-paper. Several borates are sparingly soluble, but none of them insoluble in water; whence dilute solutions of borates are non-precipitable.

a. *Alcohol* added to a mixture of sulphuric acid and a borate, burns with a marked green flame, the colour of which is best brought out on stirring the burning mixture.

β. The solution of a borate, acidified with hydrochloric acid, has the property of turning *turmeric paper* of the brown colour usually caused by alkalis. Even with dilute solutions the colouration becomes well marked on drying the paper. The acidification of not too dilute borate solutions is attended with the precipitation of boric or boracic acid H_3BO_3 .

(106.) FLUORIDES.

a. Fluorides, when gently heated with strong *sulphuric acid*, evolve hydrofluoric acid vapour HF, which becomes opaque in moist air, and has a very pungent, irritating smell. The experiment is best made in a platinum crucible, but should a test-tube or watch-glass be employed, its interior will afterwards be found corroded. When a piece of flat glass, covered with a layer of beeswax, through which some markings have been scratched, is exposed for a little while to the action of the vapour, the markings become permanently etched upon the glass, and even if very slight may be rendered evident by breathing upon its well-polished surface.

β. Fluorides, mixed with *silica* or any silicate, and warmed with *sulphuric acid*, evolve gaseous fluoride of silicon SiF_4 , as a pungent, irritating gas, rendered opaque and acid by moist air, in consequence of the following decomposition :—



Hence the wetted surface of a strip of glass, dipped into the gas, or the moist interior of a tube through which it is transmitted, as shown in fig. 38, soon acquires an opaque coating of silica.

γ. The soluble fluorides of potassium and sodium give with

chloride of calcium or *chloride of barium* a gelatinous white precipitate, soluble in hydrochloric acid, slightly soluble in ammoniacal salts, and scarcely at all soluble in acetic acid; though, indeed, the precipitated fluoride of calcium CaF_2 , or of barium BaF_2 , is often accompanied by some silicofluoride insoluble in hydrochloric acid. Fluor-spar, or native fluoride of calcium, is not readily soluble in hydrochloric acid, but an available solution may be obtained by digesting the finely-powdered spar for some time in the strong acid, and then boiling the mixture after dilution with water.

Fig. 38.

(107.) SILICATES.

Silicates of potassium and sodium, when not containing an excess of silica, are soluble in water, but all other silicates are insoluble.

a. On acidifying the solution of a silicate of alkali-metal with *hydrochloric acid* complete decomposition takes place, most usually with precipitation of some silicic acid in the gelatinous form, while another portion remains dissolved in the acid liquid. Under certain circumstances, however, as when the silicate solution is rather dilute, or when it is added at once to an excess of hydrochloric acid, there may not be any precipitation whatever. But in any case, upon evaporating down the clear acidified solution, the whole of the dissolved silicic acid separates out in the form of silica, which, after drying, is insoluble both in acid and alkaline solutions. The precipitate of gelatinous silicic acid is readily soluble in solutions of caustic alkali, and even of carbonated alkali when gently warmed therewith; but upon drying at a moderate heat, it becomes hard, gritty, and insoluble.

β. Owing to the incompetency of silicic acid to form a salt with ammonia, *chloride* or *carbonate of ammonium* added to a dissolved silicate of alkali-metal precipitates gelatinous silica,



which separates more completely on evaporation, at the same time becoming anhydrous.

γ. *Carbonate of sodium* fused before the blowpipe on a loop of platinum wire furnishes a bead which is transparent while hot, opaque when cold; but a little silica added to the strongly heated bead dissolves therein with effervescence, and, if in sufficient quantity, renders it permanently transparent.

CHAPTER III.

TOXICOLOGICAL CHEMISTRY.

(108.) For the performance of this part of the course, the student should be provided in succession with the principal poisons in the various forms in which they ordinarily occur in medico-legal practice. He must examine each poison in the several conditions in which it is presented to him, and verify all its described reactions. The different poisons may be met with in the ordinary state in which they are sold; dissolved in or diluted with water; in various organic liquids, either the vehicles in which they were administered, or the contents of the stomach for instance; mixed with solid food; in the tissues of different organs, more particularly the liver and kidney; and as stains upon clothing. They may be roughly classified into acid poisons, including the sulphuric, nitric, hydrochloric and oxalic acids; metallic poisons, including compounds of mercury, lead, copper, arsenic and antimony; and organic poisons, including prussic acid, strychnia, and opium.

§ I.—SULPHURIC ACID.

(109.) CONCENTRATED.

a. Appearance. Concentrated sulphuric acid, or oil of vitriol, occurs as a heavy, somewhat oily liquid, usually having a more or less marked brownish tint.

β. Volatility. A few drops of the acid, when cautiously

heated upon a watch-glass, or on platinum foil, disappear entirely with the formation of opaque white acrid fumes.

γ. Heat on admixture with water. Upon agitating a drachm or so of the strong acid with about an equal bulk of water, the temperature of the mixture rises very considerably, and the outside of the tube or other containing vessel becomes insupportably hot to the hand.

δ. Charring organic matter. A piece of paper, wood, or sugar, dipped into the strong acid, speedily becomes blackened or carbonised.

ε. Evolution of sulphurous acid. Upon gently heating sulphuric acid in a test tube with some chips of wood, copper turnings, or mercury, a vapour having the peculiar suffocating smell of burning sulphur is evolved; while a piece of starch-paper, moistened with iodic acid solution and held over the mouth of the tube, acquires a purple colour, which, however, the prolonged action of the vapour ultimately causes to disappear.

(110.) DILUTED.

α. Acid reaction, &c. Dilute sulphuric acid is entirely volatilisable by heat, has a marked acid reaction to test paper, and dissolves a fragment of carbonate of ammonium or sodium with rapid effervescence.

β. Charring after concentration. Marks upon paper made with the dilute acid appear simply wet, but become gradually black from carbonisation, on carefully drying the paper over a stove or gas flame.

γ. Precipitation of sulphate of barium. Solution of nitrate or chloride of barium throws down from dilute sulphuric acid a white precipitate of sulphate of barium, not affected by the addition of nitric or hydrochloric acid.

δ. Recognition of sulphur in precipitate. The precipitate having been collected on a filter, is washed, dried, and intimately mixed with about an equal quantity of black flux.

The mixture is then heated to redness in a reduction tube or before the blowpipe, whereby a fused residue is obtained, from which hydrochloric acid causes an evolution of sulphuretted hydrogen gas, recognisable by means of lead paper, to which it imparts a glistening black or brown discolouration.

(111.) IN ORGANIC LIQUIDS.

a. Sulphuric acid contained in an organic liquid, such as coffee, beer, or the contents of the stomach, readily manifests all the above described properties of the dilute acid, except that of volatility. Should the liquid be viscid or turbid, it must be diluted freely with water or proof spirit, and strained through fine muslin, or, if practicable, filtered through paper. The filtrate is precipitable by nitrate or chloride of barium (*vide* 110); and upon being evaporated down, becomes more strongly acid, chars, and evolves sulphurous acid gas (*vide* 109). Although, from the administration of antidotes in cases of sulphuric acid poisoning, the contents of the stomach or vomited matters may not exhibit any, or only a very, slight acid reaction, they may, nevertheless, yield an abundant precipitate of sulphate of barium.

(112.) STAINS ON CLOTHING.

a. The concentrated acid produces, upon black cloth, for instance, a brown stain with or without a red border; and the diluted acid a red stain, gradually becoming brown. The stains remain moist for a long time, and in all cases the fibre becomes destroyed with greater or less rapidity. By treating the stained pieces with water, a solution of sulphuric acid is obtained, which manifests acidity to test paper, and yields a white precipitate with nitrate of barium. A portion of the stained stuff heated in a reduction tube evolves sulphurous acid, recognisable by its smell, and by its reaction upon starch paper moistened with iodic acid solution.

§ II.—NITRIC ACID.

(113.) CONCENTRATED.

α. Appearance. Nitric acid when pure is colourless; but when containing peroxide of nitrogen is straw-yellow or orange, and sometimes even green or blue.

β. Volatility. The concentrated acid when exposed to the air at ordinary temperatures gives off colourless or orange fumes, having a characteristic smell, and a strongly marked acid reaction. A few drops of the acid heated upon a watch-glass disappear without leaving any residue.

γ. Action on organic matter. A strip of flannel or other nitrogenised organic tissue dipped into strong nitric acid becomes stained distinctly yellow; the colour being rendered darker and browner by the subsequent action of caustic alkali. Black and coloured cloths likewise become stained of a yellow colour, and rapidly corroded by the strong acid.

δ. Action on metals. When nitric acid is gently warmed in a test-tube with a drop or two of mercury or a few copper-turnings, violent chemical action takes place, as evidenced by the solution of the metal, and the copious evolution of orange-brown vapours which redden but do not bleach litmus paper, and produce a purple colouration on starch paper moistened with iodide of potassium.

ε. Reaction with sulphate of iron. Nitric acid added to a cold moderately concentrated solution of sulphate of iron, produces a dark greenish-brown discolouration, which disappears on the application of heat with evolution of orange fumes.

ζ. Solution of gold. Gold leaf is unacted upon by boiling nitric acid, but upon the addition thereto of a little hydrochloric acid it undergoes rapid solution.

(114.) DILUTED.

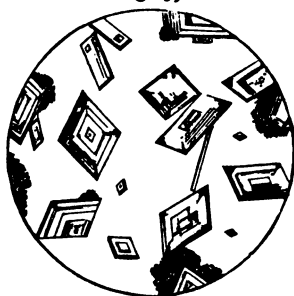
α. Acid reaction, &c. Dilute nitric acid is completely volatilised by heat; has a marked acid reaction; when evaporated from paper moistened with it, does not leave a black, but only a

slightly yellow stain; produces upon black cloth a stain, red at first, but ultimately yellow; and, if pure, gives no precipitate with solutions of chloride of barium and nitrate of silver respectively. When neutralised by the hydrate or carbonate of sodium, potassium, or barium, it is converted into a nitrate which may be obtained solid and crystalline by careful evaporation.

(115.) NITRATES.

α. Crystalline form. A drop or so of the aqueous solution of a nitrate heated slowly upon a glass plate until a solid margin appears round the edge of the liquid, yields upon cooling a well crystallised residue, which may be examined by a pocket lens or the low power of a microscope. The potassium salt crystallises in long fluted six-sided prisms, the sodium salt (fig. 39), in rhombic plates, and the barium salt in octahedrons.

Fig. 39.



β. Deflagration with charcoal. A solid nitrate heated upon charcoal, or heated with charcoal powder on platinum foil, undergoes deflagration; while a piece of filtering paper moistened with the solution of a nitrate and dried, burns in the characteristic manner of touch paper.

γ. Evolution of peroxide of nitrogen. When a mixture of a little powdered nitrate with a few copper filings is acted upon by sulphuric acid and gently warmed, orange-brown fumes are given off, which redden but do not bleach litmus paper, and produce a purple colouration on starch paper moistened with iodide of potassium. This experiment may be performed satisfactorily with less than one-tenth of a grain of nitre.

δ. Reaction with sulphate of iron. When a solution of sulphate of iron is poured carefully upon some sulphuric acid, to which a minute fragment of a nitrate has been added, or upon a

cooled mixture of sulphuric acid with a little nitrate solution, a deep greenish-brown halo is produced at the junction of the two liquids, as shown in fig. 36. This test is also extremely delicate.

ε. Solution of gold. Gold leaf when boiled in hydrochloric acid remains unchanged, but upon the addition of a little nitrate becomes dissolved wholly or in part. To demonstrate the solution of the gold in the latter case, protochloride of tin may be added, which will give rise to a purplish precipitate or turbidity.

(116.) IN ORGANIC MIXTURES.

α. Solid matters should be digested for some time in cold water, and the liquid filtered off. This method is applicable to stains on clothing, when not of too long standing. Viscid and turbid liquids are simply mixed with water and filtered. The suspected filtrate is next tested with blue litmus paper, and if found to be acid, neutralised carefully with a solution of carbonate or hydrate of sodium, evaporated down to the crystallising point, and set aside. If practicable, the resulting deposit may be collected, dried by pressure between folds of bibulous paper, dissolved in a little warm water, and the solution, filtered if necessary, evaporated, and recrystallised. Lastly, the crystalline residue is to be examined microscopically, and by the several chemical tests mentioned in the preceding section. The acid reaction may be wanting in organic mixtures, through a neutralisation of the nitric acid originally administered.

§ III.—HYDROCHLORIC ACID.

(117.) CONCENTRATED.

α. Appearance. The pure solution is colourless, or of a scarcely perceptible greenish tinge; but the commercial acid has frequently a bright yellow colour, from the presence of perchloride of iron.

β. Volatility. At ordinary temperatures the strong acid evolves colourless, almost transparent fumes, which have a marked

acid reaction, a characteristic smell, and, in common with other acid fumes, become opaque upon admixture with ammoniacal vapour. The liquid acid heated on a watch-glass evaporates without leaving any residue.

γ. Action on organic matter. Most organic tissues are gradually corroded and tinged of a yellow colour by immersion in the strong acid; but the stains produced on black cloth are at first distinctly red, and after some days reddish-brown.

δ. Want of action on metals. Hydrochloric acid boiled with a little copper or mercury simply evaporates, leaving the metal unchanged, or very nearly so.

ε. Evolution of chlorine. Peroxide of manganese warmed with hydrochloric acid in a test tube produces an abundant evolution of chlorine gas, recognisable by its greenish-yellow colour and irritating smell. It quickly bleaches litmus paper, and produces a purple colouration on starch paper moistened with iodide of potassium.

(118.) DILUTED.

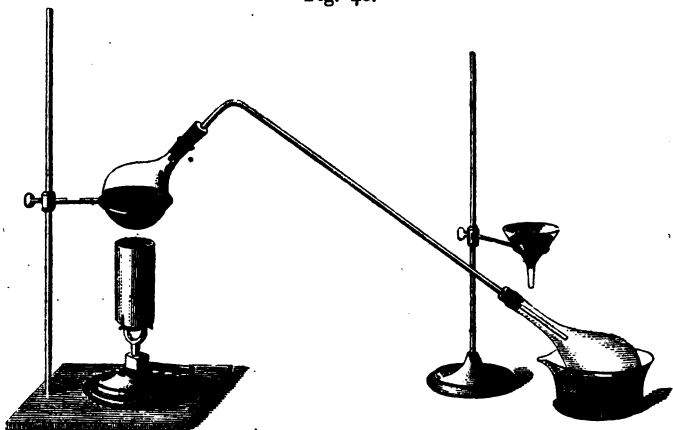
α. Neutralisation by carbonates, &c. Dilute hydrochloric acid is completely volatile, has a marked acid reaction, and dissolves most carbonates with effervescence, forming chlorides, which may be obtained in the solid state on evaporation; the alkaline chlorides, and particularly chloride of sodium, crystallising in cubical or stauroid forms. Chlorine may be liberated from the evaporated residue by the action of sulphuric acid and peroxide of manganese.

β. Precipitation of chloride of silver. Solution of nitrate of silver added to hydrochloric acid throws down a white clotty precipitate of chloride of silver, which subsides readily after brisk agitation, and by exposure to light acquires a grey or purplish colour. One portion of the precipitate may be treated with ammonia, in which it will dissolve, and another portion boiled with nitric acid, by which it will be unaffected, while the remainder may be washed, dried, and ignited in a porcelain capsule, when it will fuse into a horny mass.

(119.) IN ORGANIC LIQUIDS OR SOLIDS.

a. Distillation, &c. The liquid having shown an acid reaction to test paper, may be strained or filtered if necessary, and then distilled nearly to dryness from a retort, or a flask to which a cork and delivery tube have been adapted, as in fig. 40.

Fig. 40.



The earlier portions of the distillate are usually little else than water, but the later portions should manifest all the properties of dilute hydrochloric acid. Solid substances may be digested in distilled water, and the resulting solution examined with test paper and nitrate of silver. If possible, a portion of the liquid should be evaporated to thorough dryness, and the dissolved residue again tested with nitrate of silver to ascertain the absence or comparative absence of metallic chlorides.

§ IV.—OXALIC ACID.

(120.) SOLID.

a. Appearance, &c. Oxalic acid generally occurs in colourless, more or less well-defined four-sided prisms, which dissolve readily in boiling water to form a solution having a marked acid reaction.

β . Volatility without charring. A few crystals of the acid, when heated upon platinum foil, melt, evolve fumes, and disappear without leaving any carbonaceous residue.

γ . Effervescence with manganese. A little peroxide of manganese, free from carbonates, when added to oxalic acid moistened with water, sets up an active effervescence of carbonic acid gas.

(121.) DISSOLVED.

α . Crystalline form. One or two drops of the strongly acid liquid evaporated cautiously upon a glass plate until a solid white margin appears, yield on spontaneous cooling a crystalline residue of delicate, long flat prisms, as shown in fig 41.

Fig. 41.



β . Precipitation of oxalate of silver. Nitrate of silver produces in solution of oxalic acid, more abundantly after its neutralisation or partial neutralisation with ammonia, an opaque white precipitate of oxalate of silver, which is not discoloured by ebullition, but may dissolve if the excess of oxalic acid be large, and is readily soluble in dilute nitric acid. The precipitate collected on a filter, washed, dried, and then ignited upon platinum foil, is dissipated with a slight explosion, leaving an inconsiderable pulverulent residue of metallic silver.

γ . Precipitation of oxalate of calcium. Solution of sulphate of calcium added freely to aqueous oxalic acid, throws down a white precipitate of oxalate of calcium, insoluble in acetic, but readily soluble in dilute nitric acid. By ignition, the precipitate is converted into carbonate of calcium, which dissolves in acetic acid with effervescence.

(122.) IN ORGANIC LIQUIDS.

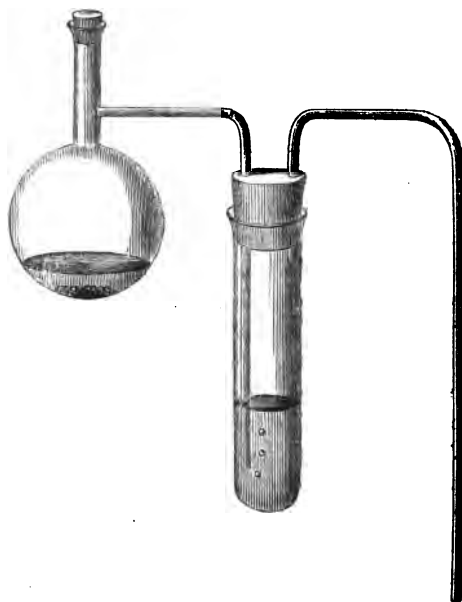
α . Reaction to test paper, &c. The acid reaction is very

decided even when the poison exists but in very small proportion. It is often necessary to filter the liquid, after previous dilution with water or proof spirit, or at any rate to strain it through muslin, before applying any reagent.

β . Precipitation of oxalate of lead. Solution of acetate of lead added to the strained or filtered liquid, throws down a precipitate of oxalate of lead, either white or discoloured by the organic matter present. The addition of the reagent is to be continued until it no longer produces a fresh precipitate. This point is easily ascertained by repeatedly testing the supernatant liquid, which separates readily after briskly agitating the mixture.

γ . Production of oxalic acid. The above precipitate of oxalate of lead having been collected on a filter and thoroughly

Fig. 42.



washed, is made into a thin magma with water, and treated with a current of washed sulphuretted hydrogen evolved from some

such arrangement as that shown in figs. 42 and 15, until the mixture, after agitation and standing at rest for a minute or so, smells strongly of the gas; when, with or without previous warming, it is to be thrown on to a filter. The filtrate will be an aqueous solution of oxalic acid, which will yield crystals on evaporation, and manifest all the above-described reactions of the dissolved acid.

δ. Production of oxalate of ammonium. Or instead of treating the washed precipitate of oxalate of lead with sulphuretted hydrogen, it may be boiled for a short time with a small quantity of dilute sulphuric acid, the mixture filtered, and the filtrate neutralised with ammonia. The resulting solution of oxalate of ammonium may be concentrated by evaporation and tested with nitrate of silver and sulphate of calcium, when characteristic precipitates of the oxalates of silver and calcium respectively will be obtained.

(123.) INSOLUBLE.

α. Oxalic acid is sometimes met with as a calcium or magnesium salt, owing to the exhibition of chalk or magnesia as an antidote. The insoluble white deposit, when boiled for some time with carbonate of sodium, yields a solution of oxalate of sodium, which after filtration and careful neutralisation with dilute nitric acid, may be tested with nitrate of silver and sulphate of calcium respectively. It is recognisable also by effervescing upon treatment with peroxide of manganese and a dilute mineral acid.

§ V.—CORROSIVE SUBLIMATE.

(124.) SOLID.

α. Appearance, solubility, &c. Corrosive sublimate usually occurs as a heavy, white, glistening powder, turned black by sulphide of ammonium, yellow by potash, and scarlet by iodide of potassium. Boiled with a little water in a test-tube it undergoes speedy solution.

β. Volatility. A small portion of the powder, heated on charcoal or platinum foil, disappears completely with production

of opaque white fumes. Heated in a narrow tube it yields a white crystalline sublimate.

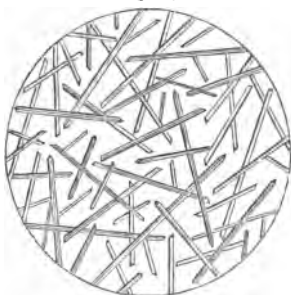
γ. Metallic reduction. When a little of the powdered salt, mixed with three or four times its bulk of recently calcined carbonate of sodium, is introduced into a thoroughly dry reduction tube, the mixture covered with some additional carbonate of sodium, and the heat of a spirit or gas-flame applied, first to the carbonate and then to the mixture, volatilisation of metallic mercury takes place, and a sublimate of well-defined mercurial globules condenses in the cold part of the tube.

δ. Detection of chlorine in residue. The bottom of the tube containing the fused residue having been snapped off at a file-mark and boiled in water, the resulting solution may be acidulated with nitric acid and tested with nitrate of silver, when a white precipitate of chloride of silver will be formed, insoluble in nitric acid, but soluble in ammonia.

(125.) DISSOLVED.

α. Crystalline form. A little of the solution evaporated cautiously on a glass plate until a solid margin appears, and set

Fig. 43.



aside to crystallise, furnishes a residue of slender opaque intersecting needles (fig. 43), which afford the above-described reactions of the solid poison.

β. Precipitation of sulphide of mercury. Sulphuretted hydrogen gas, or its aqueous solution, added to corrosive sublimate solution, produces at first a white turbidity, gradually becoming orange-brown, and finally changing into a dense black precipitate of sulphide of mercury, which may also be thrown down by sulphide of ammonium. The washed precipitate is insoluble even in boiling nitric acid.

γ. Precipitation of oxide of mercury. Excess of

potash or lime water produces an orange-yellow precipitate of mercuric oxide, the filtrate from which may be tested for chlorine by means of nitrate of silver and nitric acid.

δ. Precipitation of iodide of mercury. Iodide of potassium produces, in corrosive sublimate solution, an orange or scarlet precipitate of iodide of mercury, which disappears in excess of the precipitant, forming a colourless liquid. This reaction is very characteristic, though liable to be interfered with by the presence of various saline compounds.

ε. Reduction by a stannous salt. Protochloride of tin, added to a solution of corrosive sublimate acidulated with hydrochloric acid, produces at first a white precipitate of calomel, which, on adding more of the reagent, becomes slate-coloured, and finally almost black, from its conversion into metallic mercury. The precipitate subsides rapidly upon the application of heat; afterwards the supernatant liquid may be poured off, replaced by hydrochloric acid, and heat again applied, when the originally bulky deposit will shrink into a few globules of liquid mercury having a highly characteristic aspect.

ζ. Deposition on copper. A piece of thin copper foil or gauze immersed in the solution, previously diluted very considerably and acidulated with hydrochloric acid, quickly acquires a silver-like coating of mercury. On heating the washed and dried piece of coated copper in a subliming tube, the mercury volatilises, yielding a sublimate of metallic globules, while the copper resumes its original red colour.

η. Electrolytic test. When a few drops of the solution, preferably acidulated with hydrochloric acid, are placed on a sovereign or other piece of gold, and the coin touched through the liquid with a key or other steel instrument, a deposition of mercury takes place upon the gold at the point of contact, forming a silvery stain, which disappears upon the application of heat.

(126.) IN ORGANIC LIQUIDS AND SOLIDS.

α. General processes. The strained or filtered liquid,

acidulated with hydrochloric acid and gently warmed, may sometimes be treated with a solution of protochloride of tin, as above described; but the electrolytic test, and especially the test of metallic reduction upon copper, are in most cases much to be preferred. It is generally sufficient to prove the presence of dissolved mercury in an organic liquid, without taking the chlorine of the salt into consideration; but occasionally the entire salt may be extracted by agitating the liquid with ether, evaporating the ethereal solution, and treating the residue with water. Organic solids, thoroughly broken up or otherwise finely divided, are boiled for an hour or so with dilute hydrochloric acid, and the solution tested by immersing in it a piece of clean copper; or, if necessary, the copper may be boiled in the liquid for a considerable length of time.

β. Special process. After boiling the broken-up tissue with dilute hydrochloric acid for an hour or so as above described, and filtering off the acid decoction, the undissolved residue should be made into a thin paste with hydrochloric acid diluted with twice its bulk of water, and the mixture heated on a water bath. Then, from time to time, finely powdered chlorate of potassium is to be added little by little until the colour of the undissolved substance is reduced to a pale yellow tint, when the filtered decoction may be returned, the whole boiled for a few minutes, allowed to cool, and filtered, the filtrate evaporated to a small bulk, again filtered if necessary, and treated with excess of washed sulphuretted hydrogen gas. The resulting sulphide of mercury, mixed with much sulphur, may be collected on a filter, washed with water, dissolved in hydrochloric acid, to which a minute quantity of chlorate of potassium has been added, the liquid evaporated to dryness, the residue dissolved in water, and the resulting solution of mercuric chloride examined by any of the usual tests. This method is one of general applicability for the detection of metals absorbed into the tissues. The original precipitate produced by sulphuretted hydrogen may in all cases be dissolved in hydrochloric acid, aided by the smallest sufficient quantity of chlorate of potassium as above described, the result-

ing solution treated afresh with sulphuretted hydrogen, and the precipitate then obtained examined for mercury, lead, copper, arsenic, antimony, bismuth, tin, &c.

§ VI.—LEAD.

(127.) SOLID COMPOUNDS.

α. Metallic reduction. Lead may be readily procured in the metallic state from substances containing it in moderate quantity. When a small portion of any lead compound, mixed with three or four times its weight of carbonate of sodium, is heated on charcoal in the reducing blowpipe flame, a malleable metallic globule is soon produced, while the charcoal receives a yellow incrustation. The globule may be dissolved in dilute nitric acid, and the liquid tests applied to the solution so formed.

β. Carbonate of lead. This salt occurs as an opaque white powder, which melts and becomes yellow when heated, is turned black by sulphide of ammonium, is insoluble in water, but dissolves in dilute nitric acid with effervescence, forming a solution to which the liquid tests for lead can be applied.

γ. Acetate of lead.—Aspect, &c. Acetate of lead generally occurs as a heavy crystalline powder, of a white colour, a peculiar sour smell, and a sweetish astringent taste. It is moderately soluble in distilled water, and forms a milky liquid with common water, containing sulphates or carbonates. Its solution when evaporated upon a glass plate yields opaque white prismatic crystals.

δ. Acetate of lead.—Reactions. The salt is turned of a black colour by sulphide of ammonium, and of a yellow colour by iodide of potassium. When heated in a reduction tube, it melts, resolidifies, becomes dark in colour, gives out a smell of acetone, and leaves a carbonaceous residue containing very finely divided metallic lead. When heated in a test tube with sulphuric acid it gives off the smell of acetic acid, convertible into that of acetic ether upon the addition of a little alcohol.

Treated with solution of persulphate of iron it yields a white residue of sulphate of lead, and a dark red solution of peracetate of iron.

(128.) DISSOLVED.

a. Precipitation of sulphide of lead. Sulphuretted hydrogen gas or its solution in water, when added to any solution containing lead, gives a black or dark brown precipitate of sulphide of lead, insoluble in cold dilute hydrochloric acid, and unaffected by sulphide of ammonium. But in some cases, especially in presence of much chloride of hydrogen or iron, the precipitate comes down of a red colour, and is then turned black by sulphide of ammonium.

β. Precipitation of sulphate of lead. Dilute sulphuric acid produces an opaque white precipitate of sulphate of lead, insoluble in nitric acid, soluble in boiling and in a large excess of cold hydrochloric acid, and in a considerable excess of potash water. The precipitate is turned black by sulphuretted hydrogen or sulphide of ammonium.

γ. Precipitation of iodide of lead. Iodide of potassium added to the lead solution, which should be free from any great excess of free acid or alkali, throws down a bright yellow precipitate of iodide of lead, soluble in hot hydrochloric acid and in a large excess of potash water. It is also sparingly soluble in boiling water, and is reprecipitated on cooling in golden scales.

(129.) IN ORGANIC LIQUIDS OR SOLIDS.

a. Organic liquids. The liquid, strained or filtered if necessary, and acidulated with a few drops of nitric or hydrochloric acid, is treated with a current of washed sulphuretted hydrogen, until it acquires a marked smell of the gas, persistent after agitation. The resulting black or dark brown precipitate is allowed to subside, collected on a filter, thoroughly washed with water, and boiled, until its colour is destroyed, in nitric acid diluted with about four times its bulk of water. The solution

so obtained, filtered if necessary and concentrated by evaporation, is then submitted to the action of the ordinary tests for dissolved lead salts.

β . Organic solids. The tissue suspected to contain lead may be treated with hydrochloric acid and chlorate of potassium, as described in par. 116 β . Or it may be dried in an oven, burnt in a capsule, and the resulting charcoal heated to dull redness for several hours until a grey ash is left, which must be dissolved in dilute nitric acid by the aid of heat. The solution of the ash is then to be treated with sulphuretted hydrogen, and the black precipitate further examined.

§ VII.—COPPER.

(130.) DISSOLVED.

α . Solubility, &c. Most salts of copper are soluble in water or dilute mineral acids, except the sulphide, which, however, dissolves readily in strong nitric acid. The solutions whether aqueous or acidulous have a decidedly blue or green colour.

β . Precipitation of sulphide of copper. Sulphuretted hydrogen or an alkaline sulphide throws down a dark brown precipitate of sulphide of copper, which is scarcely affected by treatment with cold hydrochloric acid, but dissolves readily in nitric acid, is partly soluble in ordinary sulphide of ammonium, but insoluble in sulphide of sodium or potassium.

γ . Formation of cuprammonium salt. Ammonia added carefully to a cupric solution produces a bluish white precipitate, which dissolves in excess of the precipitant, forming a deep purple-blue liquid, of highly characteristic appearance, save when very dilute, in which case it resembles the similarly constituted but strong ammoniacal solution of nickel. It is, however, distinguishable therefrom by means of caustic potash, which, unless added in very large proportion, does not disturb the transparency of the cuprous, but affords a pale green precipitate with the nickel solution.

δ. **Precipitation of ferrocyanide of copper.** Ferrocyanide of potassium produces a gelatinous chocolate precipitate of ferrocyanide of copper, insoluble in mineral acids. Potash changes it into a pale blue magma readily soluble in ammonia, forming a deep blue liquid. By these properties it is distinguished from the similarly coloured precipitate of ferrocyanide of uranium. Ferrocyanide of nickel has a pale green colour.

ε. **Metallic precipitation on iron.** A steel needle or piece of polished iron wire immersed in a feebly acidulated cupric solution soon acquires a coating of metallic copper, having its characteristic red appearance. When the proportion of copper is very minute, the iron should continue immersed for several hours. Occasionally the deposit is not sufficient in quantity to present the ordinary aspect of metallic copper, but appears simply brown or black.

ζ. **Examination of metallic deposit.** The coated wire, having been washed in water, is acted upon by a drop or two of ammonia when by exposure to air the copper gradually dissolves in the ammonia, forming a deep blue solution, in which, after acidification with acetic acid, ferrocyanide of potassium causes a chocolate red turbidity of ferrocyanide of copper.

η. **Electrolytic test.** When a few drops of an acidulated solution of copper are placed in a platinum capsule, and a piece of zinc foil introduced so as to touch the capsule through the liquid, metallic copper is quickly deposited upon the platinum, either with its characteristic appearance, or simply as a brown stain. The deposit can be examined with ammonia and ferrocyanide of potassium as above described. The process possesses no advantage over that with the iron wire.

(131.) IN ORGANIC LIQUIDS AND SOLIDS.

α. **Organic liquids.** These, when containing any appreciable quantity of copper, have usually a more or less marked greenish tint. They may be acidulated with hydrochloric acid, and allowed to act upon clean iron wire as above described; or the filtered

acidulated solution may be treated with a current of sulphuretted hydrogen gas, and the resulting dark brown precipitate collected upon a filter, washed with water, and dissolved in dilute nitric or in strong hydrochloric acid. The acid solution evaporated nearly to dryness, and diluted with water, can then be examined by the usual tests.

β . Organic solids. These, when containing copper even in small quantity, acquire a deep blue colour by immersion in ammonia. They may be cut into small pieces, boiled for some time in dilute hydrochloric acid, and the resulting liquid, after being concentrated by evaporation, examined with a steel needle or a current of sulphuretted hydrogen gas (*a*). Or the tissue may be destroyed, either with hydrochloric acid and chlorate of potassium, as described in par. 116 β , or by incineration, as described in par. 119 β , and the residues further examined.

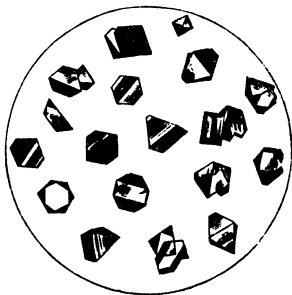
§ VIII.—ARSENIC.

(132.) ARSENIOS ACID OR ANHYDRIDE.

a. Appearance. Commercial arsenious acid, white arsenic, or arsenious anhydride, usually occurs as a heavy white powder, but may be met with in transparent vitreous masses, or in opaque porcellaneous masses, or in masses which are opaque externally and glassy in the centre.

β . Volatility. A minute quantity of the powder heated on platinum foil volatilises entirely with evolution of opaque white fumes; any fixed residue being due to impurity, probably sulphate of calcium. A little of the powder heated in a subliming tube also volatilises, and deposits an iridescent sublimate in the cool part of the tube. Or the volatilisation may be effected in a short wide test tube, and the sublimate condensed on a flat strip of glass held over the

Fig. 44.



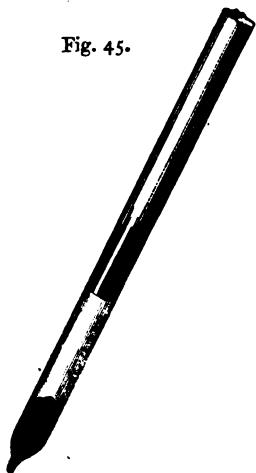
mouth of the tube. Upon examination with a lens, the sublimate in the tube or on the glass will be seen to consist chiefly of octahedral crystals, as shown in fig. 44, which do not polarize.

γ. Action of sulphide of ammonium. When a drop or so of sulphide of ammonium is added to a little white arsenic contained in a watch glass, there is no alteration of colour produced, but on applying a gentle heat, solution takes place, and on evaporating to dryness, a yellow film of sulphide of arsenic is left, soluble in alkalies, insoluble in hydrochloric acid.

δ. Solubility. Powdered white arsenic when agitated with water in a test tube, does not perceptibly dissolve, but remains partly as a film over the surface, partly in small aggregations at the bottom. This state of immiscibility, which is very characteristic, does not disappear even on prolonged boiling. By filtration, however, a clear aqueous solution of arsenious acid is obtained. Moreover on adding a little potash or hydrochloric acid to the hot mixture of water and white arsenic, complete solution is very easily effected.

ε. Reducibility. A minute quantity of the powder sprinkled upon red-hot charcoal evolves scarcely visible vapours having a peculiar garlic-like odour. When a little white arsenic mixed with three or four times its bulk of dry soda-flux, produced by the incineration of acetate of sodium with some additional charcoal, is introduced into a narrow reduction tube of hard glass, made perfectly dry and warm, there is produced, on subjecting the mixture to the heat of a spirit flame, a sublimate of reduced arsenic which condenses in the upper cool part of the tube in the form of a metallic ring, as shown of the actual size in fig. 45.

Fig. 45.



Or a minute quantity of white arsenic may be placed in the reduction-tube, covered with a considerable thickness of powdered charcoal, and heat applied from above to below, so that the arsenic may volatilise through the red-hot charcoal, when a ring of reduced metal will appear as before.

ζ. Characters of the metallic ring. The arsenical ring is characterised as follows:—*a.* By its lustrous steel grey appearance. Should it present an opaque brownish-black colour, its proper aspect may be brought out by the cautious application of heat, when the characteristic metallic grey ring will remain, and a more volatile dark-coloured compound of arsenic be volatilised. The interior surface of the metallic sublimate, rendered visible by breaking the tube, presents a crystalline appearance. *b.* By its volatility. Upon heating the sublimate to a temperature considerably below redness, it may be readily volatilised from one part of the tube to another. *c.* By its conversion into arsenious anhydride. After repeated volatilisations up and down the tube, the ring of metal is gradually replaced by a ring of iridescent crystals of white arsenic, shown by a lens to consist of variously modified octahedrons. These may be boiled in a small quantity of water for some time, when a solution of arsenious acid will be formed, to which the ordinary liquid reagents can be applied. *d.* By its conversion into arsenic acid. The ring of metal, when warmed with a drop or two of nitro-muriatic acid, disappears, and on evaporating to dryness, a residue of arsenic acid is left, which may be dissolved in water, and tested with nitrate of silver solution, when a brick-red precipitate of arseniate of silver will be produced.

(133.) DISSOLVED.

a. Reaction, &c. The aqueous solution of arsenious acid is clear, colourless, tasteless, inodorous, and has a faintly acid reaction to test-paper. When evaporated upon a glass plate, it leaves a white residue of minute octahedral non-polarising crystals, which may be volatilised by a further application of heat.

β. Precipitation of arsenite of silver. Ammonio-nitrate of silver, added to aqueous arsenious acid, throws down an opaque yellow precipitate of arsenite of silver, soluble in ammonia and in dilute nitric acid. The reagent is made by adding a weak solution of ammonia drop by drop to a strong solution of nitrate of silver, until the brown precipitate at first produced is just redissolved.

γ. Precipitation of arsenite of copper. Ammonio-sulphate of copper, added to aqueous arsenious acid, produces a light green precipitate of arsenite of copper, soluble in ammonia and in dilute acids. When collected on a filter, washed, dried, and heated in a reduction tube, it yields a crystalline sublimate of arsenious anhydride. Ammonio-sulphate of copper is made by carefully adding ammonia drop by drop to a somewhat dilute solution of sulphate of copper, until the precipitate at first produced is nearly redissolved.

δ. Precipitation of trisulphide of arsenic. When a current of washed sulphuretted hydrogen gas is passed through a solution of arsenious acid acidulated with hydrochloric acid, an abundant bright yellow precipitate of arsenious sulphide or orpiment is produced. Sulphide of ammonium does not give any precipitate with aqueous arsenious acid until some other acid is added; while sulphuretted hydrogen produces only a yellow discolouration.

ε. Characters of the above precipitate. The yellow precipitate may be collected on a filter, washed with water, and tested as follows. Boiled with hydrochloric acid it does not perceptibly dissolve, but, on the addition thereto of a little nitric acid, disappears with the formation of red fumes, a globule of melted sulphur frequently remaining undissolved. Treated with sulphide, hydrate, or carbonate of ammonium, it undergoes solution, and is again thrown down by the addition of hydrochloric acid. Dried and heated in a reduction tube, it sublimes unchanged or nearly so. Dried and heated in a reduction tube, after thorough mixture with four or five times its bulk of soda-flux, it yields a sublimate or ring of reduced metal.

(134.) MARSH'S TEST.

a. Nature of process. When a substance containing arsenic is acted upon by nascent hydrogen, usually developed by the reaction of metallic zinc and dilute sulphuric or hydrochloric acid, a gaseous compound of arsenic and hydrogen, known by the name of arsenetted hydrogen, is given off. This gas may be generated in any ordinary form of hydrogen-apparatus; even the adaptation to one another of a flat-bottomed vial, perforated cork, and piece of glass tube, either straight or bent, according to circumstances, and drawn out to a moderately fine point, will answer the purpose. The arrangement shown in fig. 47 is in some cases very convenient; while in others, the original bent tube devised by Marsh may be most advantageously employed. It may be made quite plain, as in fig. 46, or be provided with a couple of large strong bulbs, one in the upper part of the long, and one in the lower part of the short limb. This short limb is furnished with a movable stopcock, into which is screwed either a short jet, for burning the issuing gas, as shown in the figure, or an elbow, to which a horizontal piece of glass tubing may be readily adapted. Arsenetted



hydrogen is identified by its property of yielding deposits of metallic arsenic, either upon imperfect combustion, as originally pointed out by Marsh, or by exposure to a dull red heat, as recommended by a committee of the French Academy.

(135.) ORIGINAL MARSH'S PROCESS.

When it is intended to obtain deposits by an imperfect combustion of the gas, Marsh's apparatus (fig. 46) is usually employed in the following manner. The stopcock being removed, a piece of stout glass rod is carefully dropped into the shorter limb of the tube. It should be sufficiently small to reach the bend, but not small enough to pass into the longer limb: a glass stopper will often answer the purpose extremely well. Two or three compact lumps of metallic zinc are then let fall upon the piece of glass, the open stopcock replaced, and cold diluted sulphuric acid, in the proportion of about one part of acid to six or seven of water, poured into the longer limb; so that when the liquid is level in the two limbs, there may yet remain some little free space beneath the stopcock. The dilute acid is allowed to act upon the zinc for a few minutes, and the stopcock then closed, whereby the shorter limb becomes gradually filled with hydrogen gas, the acid being gradually driven up into the longer limb. The stopcock is then opened, and the issuing hydrogen quickly inflamed. It ought to burn with a scarcely visible flame, which should not produce a deposit or even a discolouration upon a piece of clean glass or porcelain momentarily depressed upon it. As soon as all the hydrogen is driven out of the shorter limb by the descending acid, the stopcock is reclosed while another accumulation of gas takes place, which is then released, inflamed, and examined as before. Or the issuing gas, without being inflamed, may be allowed to impinge on paper moistened with nitrate of silver solution, which should not thereby acquire any discolouration. When, after several examinations, the purity of the hydrogen, and consequently of the materials used to generate it, has been satisfactorily ascertained, the arsenical liquid is introduced and the experiment repeated.

a. Appearance of flame, &c. The hydrogen gas subsequently evolved is contaminated to a greater or less extent with arsenetted hydrogen, and produces a metallic-looking discolouration upon paper moistened with nitrate of silver solution. It

burns with a bluish flame, and evolves a white smoke of arsenious anhydride.

β. Reactions of the smoke. The smoke or vapour may be tested by holding over the summit of the flame a piece of porcelain moistened with ammonio-nitrate of silver, when a yellow turbidity due to arsenite of silver will make its appearance; or by enclosing the flame more than once if necessary by a short wide test-tube moistened on its interior with water, whereby a weak solution of arsenious acid will be formed, which may be tested with sulphuretted hydrogen, or other appropriate reagent.

γ. Formation of deposit. When a piece of clean glass, porcelain, or talc is momentarily depressed upon the flame so as to cut off about two-thirds of its height, there is produced upon the cold surface a dark stain or deposit which is constituted of three products, in the centre of metallic arsenic, in the exterior of arsenious anhydride, and in the intermediate zone of a compound considered to be a suboxide of arsenic. That the stain is really arsenical is shown by its possessing the following characters:—

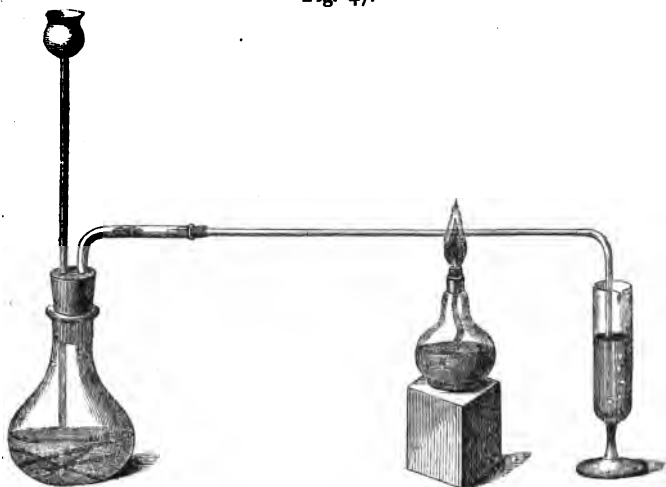
a. Metallic brilliancy. The lustrous appearance of the arsenical stain is best seen on its free surface, but is recognisable through the glass. **b. Hair-brown colour.** This colour appertains particularly to slight stains, and to the intermediate portion of larger stains, in which last it is best manifested by means of transmitted light. **c. Volatility.** The arsenical stain disappears readily on the application of a heat considerably below redness; this property of volatility is very evident when the stain has been produced on a thin plate of talc. During its volatilisation the metallic arsenic becomes converted into arsenious anhydride. **d. Solubility in chloride of lime.** When the arsenical stain is warmed with a few drops of bleaching liquid, complete solution speedily takes place. For the application of this test it is convenient to produce the stain on the interior of a watch-glass. **e. Non-solubility in cold disulphide of ammonium.** The arsenical stain is not perceptibly affected by treatment with a drop of yellow sulphide of ammonium solution. But on heating to dryness, a bright yellow stain of orpiment is

produced, containing a dark nucleus of undissolved metallic arsenic. (Guy.) *f.* Conversion into arsenic acid. The arsenical stain disappears completely when gently warmed with a drop or two of nitric or nitro-muriatic acid. By evaporating to dryness, a slight residue of arsenic acid is left, recognisable by its ready solubility in a drop of water, so as to form a solution in which nitrate of silver produces a brick-dust-red turbidity.

(136.) MODIFIED MARSH'S PROCESS.

Marsh's apparatus may also be used for generating arsenetted hydrogen when it is intended to decompose the gas by extraneous heat. In this case the jet of the stopcock is unscrewed and replaced by a metal elbow, to which a horizontal piece of narrow hard glass tubing is adapted. But some such apparatus as that shown in fig. 47 is on the whole to be preferred. It consists of a

Fig. 47.



small flask furnished with a cork through which passes a long funnel tube, and a short wide rectangular tube, loosely plugged with a little cotton wool, and connected by means of a perforated

cork with a long horizontal piece of narrow hard glass-tubing, bent downwards at its extremity so as to dip into a solution of nitrate of silver. The apparatus is charged with pure zinc and dilute sulphuric acid, so as to cause an evolution of hydrogen gas, which passes through the cotton wool, where it deposits any mechanical impurities, along the horizontal tube, and into the silver solution. During the transmission of the gas, the flame of a spirit lamp is applied steadily to some particular part of the horizontal tube, and if after a little time there is not produced any deposit within the tube, or precipitate in the silver solution, the materials are known to be pure, and the arsenical liquid may be introduced.

a. Decomposition by heat. The resulting arsenetted hydrogen is decomposed in its passage through the heated portion of the tube, and deposits a steel-grey ring of metallic arsenic at some little distance beyond the flame. The ring may be identified by its appearance, its position at a little distance beyond the flame, its volatility, its conversion into arsenious anhydride by repeated sublimations, and its conversion into arsenic acid by treatment with nitro-muriatic acid and evaporation to dryness.

β. Decomposition by nitrate of silver. Any arsenetted hydrogen that may escape decomposition by the flame, or that may be purposely allowed to escape, is arrested by the nitrate of silver solution, with formation of a black deposit of metallic silver. On the termination of the experiment, the excess of silver may be precipitated with hydrochloric acid, the filtrate evaporated to dryness, and the residue of arsenic acid dissolved in water and tested with the usual reagents.

(137.) REINSCH'S TEST.

This test is particularly useful for the detection of arsenic in organic liquids or solids. The suspected liquid is simply acidulated with about one-eighth of its bulk of pure hydrochloric acid, and boiled. The solid tissue is cut up into very small pieces and boiled for some time in a mixture of about one part of hydrochloric acid with six of water.

α. Deposit on copper. A small piece of clean copper foil, or preferably of fine gauze, is introduced into the hot acidulated liquid, and the boiling continued for a period varying from a few minutes to a quarter of an hour or longer. Should the copper acquire a grey metallic discolouration, other pieces thereof may be added from time to time, and the supply continued so long as the last added piece assumes any perceptible alteration in colour.

β. Character of deposit. The pieces of coated foil or gauze are removed from the liquid, washed in water, and dried between folds of bibulous paper. The deposit unless very thick adheres firmly to the subjacent copper, presents a well-marked metallic lustre, and has a dark steel-grey colour, or, if very thin, a somewhat bluish tint. On the application of heat it disappears entirely, while the copper resumes its ordinary appearance.

γ. Crystalline sublimate. A piece of the coated copper, held between the fingers, is warmed over a flame, coiled up into a small bulk, and introduced into an ordinary reduction tube (fig. 45). The heat of a small spirit flame is then carefully applied, at first a little above the coil, and afterwards to the coil itself, whereby the arsenic is volatilised, oxidised, and condensed in the cool part of the tube as a crystalline sublimate. If necessary, several pieces of coated copper may be thus heated successively in the same tube until a sufficiently obvious sublimate of arsenious anhydride is produced, which, when examined by a lens or the low power of a microscope, will exhibit highly iridescent octahedral forms. A small piece of tubing open at both ends, one of which is drawn out to a long, almost capillary, termination, as

Fig. 48.



shown in fig. 48, is convenient for volatilising a very slight deposit. The coated foil having been introduced and tilted down

to the shoulder, the tube is sealed by the blowpipe at the point *a*, and the resulting cylindrical bulb containing the foil heated in a spirit-flame, from its capillary shoulder backwards to its sealed extremity, whereby a crystalline ring becomes condensed in the capillary projection at *b*.

δ. Reactions of the sublimate. By means of a couple of file-marks, the short length of tubing containing the sublimate may be broken off from the two ends of the tube, and the sublimate itself be acted upon by reagents. Moistened with sulphide of ammonium solution and dried in a water-bath, it yields a yellow residue of orpiment. Moistened with a mixture of nitric and a little hydrochloric acid and evaporated to dryness, it leaves a slight residue of arsenic acid, which produces a red turbidity when treated with a drop of nitrate of silver solution.

(138.) IMPEDIMENTS TO REINSCH'S TEST.

α. Influence of oxygenants. Reinsch's process is not applicable in the presence of oxidising bodies, which moreover enable dilute hydrochloric acid to dissolve metallic copper. But the majority of such compounds may be reduced by the action of sulphite of sodium upon the acidified liquid, while any excess of sulphurous acid from the decomposition of the salt may be got rid of by ebullition before introducing the foil or gauze.

β. Purity of the acid. This may be ascertained by diluting a sufficient quantity of the hydrochloric acid with about four times its bulk of water, and boiling a very small piece of foil or gauze in the diluted liquid for a period of twenty minutes or half an hour.

γ. Purity of the copper. As even in the most satisfactory performance of Reinsch's test, there is always some, although but an extremely minute quantity of the copper dissolved, and as commercial copper is rarely quite free from arsenic, it is important that the foil or gauze employed in the experiment should be specially tested as to its purity. If, however, the

solution of four or five grains of the copper does not yield any evidence of arsenic, the metal is quite pure enough for the purpose, even though a trace of arsenic should be detected in a larger quantity of it. A few grains of the copper cut into fine pieces are placed in a small tube-retort, or in a bulb-tube, such as that shown in fig. 14, with not less than twice their weight of precipitated peroxide of iron, and an excess of hydrochloric acid. The mixture is then distilled to dryness, great care being taken at the last to prevent spurting. Any arsenic originally contained in the copper is in this manner carried over in the form of chloride of arsenic, and may be condensed in a little water with the excess of aqueous hydrochloric acid. The resulting liquid may then be tested for the presence of arsenic by boiling in it a fresh piece of clean copper gauze or foil.

δ. *Modified processes.* The peroxide of iron mentioned above may be replaced by an equivalent quantity of perchloride of iron. Indeed it is better to dissolve the peroxide in excess of hydrochloric acid, and then employ the residue left on evaporating to dryness, which will be free from any trace of arsenic the peroxide itself may have originally contained. Moreover, oxide or chloride of copper may be substituted for the peroxide or perchloride of iron, though not with advantage. Or the copper may be dissolved in hydrochloric acid alone, without the addition of any special oxygenant, by moistening the metal with the acid and exposing both to the air for several days. The addition of a few drops of a solution of perchloride of iron or chloride of copper to the acid greatly facilitates this solution by exposure to air. The hydrochloric solution, no matter how obtained, is eventually distilled to dryness, and the distillate tested for arsenic.

(139.) OTHER FORMS OF ARSENIC.

a. *Orpiment and realgar.* These sulphides of arsenic are yellow or orange-coloured compounds, which volatilise unchanged upon the application of heat. Mixed with soda flux and heated in reduction tubes, they give rise to sublimes of metallic

arsenic; and the residues, when moistened with hydrochloric acid, evolve sulphuretted hydrogen, recognisable by its smell and action on lead paper. Orpiment and realgar are not dissolved by boiling hydrochloric acid, but disappear more or less completely in nitro-hydrochloric acid, forming solutions from which arsenic acid may be obtained by evaporating to dryness. They also dissolve in sulphide of ammonium, and are redeposited on evaporating the liquid to dryness.

β. Scheele's green. This well-known green pigment is an impure arsenite of copper. Heated in a reduction tube, it yields a crystalline sublimate of arsenious anhydride, and a black residue of oxide of copper, which may be dissolved in hydrochloric acid, and tested by the usual reagents. Arsenite of copper dissolves in dilute hydrochloric acid, forming a solution in which, after precipitation of the copper by excess of oxalate of ammonium, arsenic may be detected by sulphuretted hydrogen, or by Marsh's or Reinsch's tests.

(140.) IN ORGANIC MIXTURES.

a. A hydrochloric acid decoction may be prepared as already described and tested by Reinsch's process. Or the organic substance may be distilled to dryness with hydrochloric acid, the residue redistilled with fresh hydrochloric acid, the two distillates collected in water, and the product examined by sulphuretted hydrogen, or by Marsh's or Reinsch's process. Or the tissue, &c., may be destroyed with hydrochloric acid and chlorate of potassium, the solution submitted to the prolonged action of sulphuretted hydrogen, and the resulting precipitate further examined (vide par. 116 β).

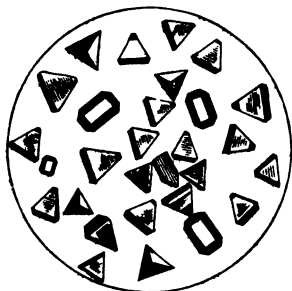
§ IX. ANTIMONY.

(141.) ANTIMONIAL SALTS.

a. Tartar emetic. This compound usually occurs as a white powder, or in ill-defined crystalline masses. It becomes charred by heat, and when ignited with a little carbonate of sodium on charcoal before the blowpipe, yields a bead of brittle metal and

an abundant white incrustation. It is turned of an orange colour, and finally dissolved by sulphide of ammonium. Its solution in

Fig. 49.



water when carefully evaporated yields beautifully polarising crystals (fig. 49), chiefly tetrahedral, but here and there cubical and octahedral. When acidified by nitric or hydrochloric acid, it furnishes a white precipitate soluble in excess of either acid.

β . Chloride of antimony. This is a highly corrosive fuming liquid, usually having a yellow or orange colour from the presence of chloride of iron. Poured into water, it gives rise to an abundant white precipitate, which, after thorough washing with water, is turned of an orange colour and finally dissolved by sulphide of ammonium. The precipitate is also readily soluble in tartaric acid. Mixed with carbonate of sodium and heated on charcoal before the blowpipe, it yields a bead of brittle metal and an abundant white incrustation.

(142.) IN SOLUTION.

α . Precipitation of trisulphide of antimony. A current of sulphuretted hydrogen gas passed into an antimonial solution acidified with tartaric acid, throws down an orange precipitate, which after washing with water, is insoluble in carbonate, but soluble in sulphide of ammonium, forming a solution from which it may be reprecipitated on the addition of an acid.

β . Precipitation of oxichloride of antimony. Precipitated trisulphide of antimony dissolves completely in hot hydrochloric acid with evolution of sulphuretted hydrogen gas. The resulting trichloride of antimony, freed from excess of hydrochloric acid by evaporation to a small bulk, gives when poured into water an abundant white precipitate of oxichloride of antimony, soluble in tartaric acid.

γ Metallic precipitation on tin. A piece of tin foil or bar immersed in the above tartaric acid solution, becomes speedily covered with a pulverulent black deposit of metallic antimony.

(143.) **MARSH'S TEST.**

The antimonial solution, acidulated with tartaric acid, may be introduced into the original or modified form of Marsh's apparatus, previously charged with pure zinc and dilute sulphuric acid, and the resulting gas examined as follows:—

α. Appearance of flame, &c. Hydrogen contaminated with antimonetted hydrogen produces a black discolouration on paper moistened with nitrate of silver solution. It burns with an opaque bluish white flame, and evolves a white smoke of teroxide of antimony, which, unlike the arsenical smoke, does not produce a yellow turbidity with ammonio-nitrate of silver.

β. Characters of deposit. When a piece of talc, porcelain, or glass is depressed upon the flame, a dark stain or deposit is produced, distinguishable from the arsenical stain by the following properties:—*a*, by its comparative want of metallic lustre; *b*, by its smoky black colour; *c*, by its non-volatility save at a heat approaching redness; *d*, by its insolubility in chloride of lime; *e*, by its ready solubility in yellow sulphide of ammonium, so as to form a solution which on evaporation to dryness leaves a bright orange stain; and *f*, by its yielding after treatment with nitro-muriatic acid and evaporation to dryness, a residue which does not give a red precipitate with nitrate of silver solution.

γ. Decomposition by heat. Instead of burning the antimonetted hydrogen, it may be transmitted through a tube heated to redness, and finally through a solution of nitrate of silver. The deposit of antimony produced in the tube is characterised by its position, just before and beyond the exact spot where the heat is applied, by its want of volatility, by its non-

convertibility into arsenious anhydride or arsenic acid, and by its ready solubility in yellow sulphide of ammonium to form a solution which leaves a bright orange stain on evaporation.

δ. Reaction with nitrate of silver. Antimonetted hydrogen produces in nitrate of silver solution a black deposit of antimonide of silver, from which, after washing with water, the antimony may be dissolved away by a boiling solution of cream of tartar, and precipitated from the resulting solution by sulphuretted hydrogen.

(144.) REINSCH'S TEST.

α. Deposit on copper. The deposition of antimony upon copper foil or gauze boiled in a hydrochloric acid decoction of organic matter contaminated with antimony, or in a weak acidulated solution of some antimonial salt, takes place exactly as does the deposition of arsenic under similar circumstances. The highly lustrous deposit of antimony differs from that of arsenic in having a marked violet colour, and in being less easily dissipated by heat. When a piece of the coated foil or gauze is strongly heated in a reduction tube, it either does not afford any sublimate at all, or else a very slight white deposit situated close to the heated end of the tube, not having a crystalline character, and being practically non-volatile.

β. Solution of deposit. When boiled for a few minutes in a weak feebly alkaline solution of permanganate of potassium, the antimonial coating is dissolved away from the copper, while the permanganate loses its colour and furnishes a slight turbidity of manganic hydrate. The filtered liquid acidulated with hydrochloric acid and treated with sulphuretted hydrogen, acquires a yellowish colour, and on standing deposits an orange precipitate, which may be further examined if necessary. Or, if the antimonial coating be not very thick, it will suffice to boil the copper for some time, with frequent exposure of its surface to the air, in a weak solution of caustic potash only, and to treat the resulting liquid, after acidification by hydrochloric acid, with sulphuretted hydrogen.

(145.) IN ORGANIC MIXTURES.

The processes of precipitation by sulphuretted hydrogen and deposition on copper, are perfectly applicable to acidified organic liquids, and to the hydrochloric acid decoctions of organic tissues. Or the tissue may be destroyed by hydrochloric acid and chlorate of potassium, the solution after evaporation treated by sulphuretted hydrogen, the resulting precipitate dissolved in boiling hydrochloric acid, and the solution so formed tested in Marsh's apparatus, or by the process of Reinsch.

§ X.—PRUSSIC ACID.

(146.) IN AQUEOUS SOLUTION.

a. Appearance, &c. Prussic or hydrocyanic acid CNH , occurs in the state of aqueous solution as a colourless, perfectly volatile, feebly acid, mobile liquid. Its vapour, which is given off at all ordinary temperatures, is invisible, has an odour said to be like that of bitter almonds, and when inspired even in minute quantity causes a peculiar sensation in the fauces.

β. Formation of Prussian blue. The turbid greenish liquid made by adding excess of potash to solution of ordinary sulphate of iron, does not undergo any visible alteration when mixed with aqueous prussic acid; but, on acidifying the mixture with hydrochloric acid, a bright blue, or sometimes a greenish blue colour is developed, due to the production of finely divided Prussian blue, which gradually separates as a distinct precipitate. Or the potash and sulphate of iron may be added separately to the suspected liquid, and the mixture be afterwards acidulated with hydrochloric acid.

γ. Formation of sulphocyanate of iron. Aqueous prussic acid, mixed with a drop or two of yellow sulphide of ammonium solution and evaporated to dryness at a low temperature, leaves a residue of sulphocyanate of ammonium, which, when moistened with water and tested with a drop of per-

chloride of iron, produces a dark-red solution of persulphocyanate of iron.

δ. Formation of cyanide of silver. Solution of nitrate of silver added to aqueous prussic acid throws down a white precipitate of cyanide of silver, which quickly subsides after agitation. It is not affected by cold nitric acid, but when separated from the supernatant liquid dissolves more or less completely in the strong boiling acid. The precipitate collected on a filter, washed, dried, and heated in a reduction tube, evolves cyanogen gas, which if ignited at the mouth of the tube, will burn with its peculiar rose-coloured flame.

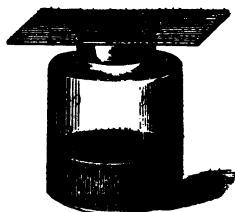
ε. Decompositions of precipitate. The precipitate treated with hydrochloric acid evolves prussic acid vapour, which may be received on the interior of a watch-glass moistened either with yellow sulphide of ammonium, or with a mixture of potash and sulphate of iron, as described under the head of the vapour reactions. Or a portion of the precipitate may be treated with a drop of yellow sulphide of ammonium, dried at a low temperature, and the residue, after being moistened with water, tested by a drop of perchloride of iron, when a dark red liquid will be produced, easily distinguishable from the black precipitate of sulphide of silver. Or a portion of the precipitate may be treated first with potash, then with a drop of sulphate of iron, and lastly with a little hydrochloric acid, when Prussian blue will be formed, together with white chloride of silver.

(146.) IN VAPOUROUS STATE.

The succeeding tests may be applied to the vapour of the pure liquid acid, or to the vapour produced by the action of hydrochloric acid upon precipitated cyanide of silver, or to the vapour evolved spontaneously from organic liquids or solids containing prussic acid. Organic substances which do not react satisfactorily with these vapour tests may be distilled in a water-bath, and the distillate treated similarly to the pure aqueous acid as above described.

a. Formation of Prussian blue. When a mixture of potash and sulphate of iron, smeared upon the interior of a watch-glass, or preferably on a flat glass slip (fig. 50), is exposed for a few minutes to the action of prussic acid vapour, there is produced on acidification with hydrochloric acid, a solution of the iron magma and development of Prussian blue.

Fig. 50.



β. Formation of sulphocyanate of iron. A drop of yellow sulphide of ammonium placed on a watch-glass or glass slip and exposed for a short time to the action of prussic acid vapour, yields, when evaporated to dryness at a low temperature, a residue of sulphocyanate of ammonium, which produces a dark red colour, on the addition of perchloride of iron.

γ. Formation of cyanide of silver. A drop of nitrate of silver, placed on a watch-glass or glass slip and exposed to the action of the vapour, becomes white and opaque from the formation of cyanide of silver, convertible into Prussian blue, or sulphocyanate of iron, as previously described.

When the prussic acid vapour from some organic mixture is contaminated with sulphuretted hydrogen, it produces a blackening of the silver salt; but no interference with the sulphocyanate reaction is manifested under the same circumstances.

§ XL.—STRYCHNIA.

(148.) IN PURE STATE.

a. Nature, solubility, &c. Strychnia is a vegetable alkaloid, having the formula $C_{27}H_{45}N_3O_4$. It is more or less freely soluble in alcohol, chloroform, benzole, and ether; scarcely at all soluble in pure water; but readily soluble in acidulated water. It is capable of uniting with and neutralising acids, to form definite crystallisable salts, of which the sulphate, nitrate, hydro-

chlorate, oxalate, tartrate, and acetate are soluble in water. Most other strychnia compounds are more or less insoluble, whence solutions of strychnia salts are precipitated by a very great number of reagents, including hydrate, carbonate, iodide, sulphocyanate and chromate of potassium, carbazotic acid, phospho-molybdate of sodium, iodide of potassium with iodine, potash double iodide of mercury and potassium, perchloride of platinum, trichloride of gold, &c.

β. Appearance. Strychnia usually occurs in the form of a crystalline powder, or of well-defined prismatic crystals, either white, or of a pale buff colour. The ordinary salts of strychnia are generally met with as crystalline powders. Strychnia and its salts when heated, melt, burn with a smoky flame, and leave a carbonaceous residue.

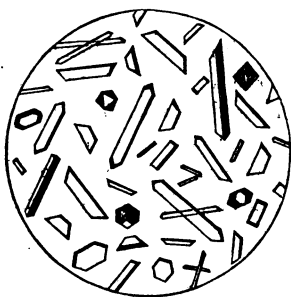
γ. Bitter taste. The bitterness of strychnia is peculiar, and has been sometimes spoken of as metallic. Its intensity is so great, that one drop of a gallon of water, in which a grain of strychnia is dissolved, presents a recognisable bitter taste; while with $\frac{1}{10000}$ part of strychnia in solution, the bitterness is well marked and persistent. The taste of strychnia salts is but slightly less intense than that of the alkaloid itself. In very dilute solutions only is the bitterness capable of partial concealment by other sapid bodies.

δ. Crystalline form. A drop or so of a spirituous or ethereal solution of strychnia allowed to evaporate spontaneously on a glass slip, furnishes a crystalline residue consisting of rectangular prisms often terminated by double or single oblique planes, and in variously modified octahedra, as shown in fig. 51. The forms deposited from a chloroformic solution are, for the most part, not well characterised.

ε. Precipitation. Aqueous solutions of strychnia salts, containing from $\frac{1}{1000}$ to $\frac{1}{500}$ part of strychnia, are precipitated by the several reagents mentioned above, either immediately or on stirring; the most delicate though least characteristic precipitants being the phospho-molybdate of sodium, and the potash.

solution of hydrargyro-iodide of potassium. Hydrate or carbonate of potassium causes a gradual deposition of well-defined strychnia crystals, insoluble in excess of the precipitant (fig. 51). The precipitates thrown down by iodide, sulphocyanate, and chromate of potassium, carbazotic acid, and the chlorides of platinum and gold, are also crystalline.

Fig. 51.



ζ. Action of acids. Strong sulphuric acid is without action on strychnia, even at and above the temperature of boiling water. Strong nitric acid usually produces a yellow, or yellow-brown, discolouration; but is said to be without visible action on perfectly pure strychnia, although this seems doubtful.

η. Colour tests. When a little peroxide of lead is added to a fragment of strychnia, dissolved in a drop of strong sulphuric acid mixed with $\frac{1}{4}$ of its bulk of strong nitric acid; or preferably, when a little peroxide of manganese, or bichromate, or ferridcyanide, or permanganate of potassium, is added to a fragment of strychnia dissolved in a drop of strong sulphuric acid, there is produced a magnificent purple-blue colour, becoming gradually crimson, and finally reddish pink. The delicacy of this test, when special precautions are taken, is almost illimitable, less than $\frac{1}{100000}$ of a grain having been stated to give the reaction. With from $\frac{1}{5000}$ to $\frac{1}{10000}$ of a grain it is easily obtainable. In operating on small quantities, the following plan may be adopted with advantage. The dry strychnia, usually the residue of an evaporation, in which case it must be allowed to become quite cold, is moistened with the smallest sufficient quantity of strong sulphuric acid. By the side of it is next placed a minute drop of a mixture of sulphuric acid with a little very finely-powdered amorphous peroxide of manganese, and the two then brought into contact. The experiment should be made on a surface of

white porcelain, or on a flat watch-glass or glass slip, resting on a sheet of white paper.

θ. Physiological test. When a minute quantity of solid or dissolved strychnia is introduced underneath the incised skin of a small frog, well-marked tetanic convulsions are manifested by the animal, usually within a quarter of an hour; and with a strong dose almost immediately. This tetanus is said to have been produced with so small a quantity as $\frac{1}{5000}$ of a grain of strychnia; but the delicacy of the test varies much with the state of the animal, freshly-caught young frogs being the most excitable.

(149.) IN ORGANIC MIXTURES.

α. If a liquid, it is merely acidified, mixed in some cases with a little spirit of wine, filtered, evaporated nearly to dryness, and the residue extracted with strong alcohol. If a solid, it is brought into a state of fine division, and mixed with a little proof spirit acidulated with dilute sulphuric or other acid, acetic, oxalic, tartaric, &c. After digestion for some time in a water bath, the mixture is filtered, the insoluble matters washed with proof spirit, the washings added to the filtrate, the whole of the clear liquid evaporated down to a small bulk, and the residue so obtained extracted with strong alcohol. The alcoholic solution is then evaporated to dryness, the residue dissolved in a little water, the liquid filtered into a long tube or bottle, and rendered alkaline with carbonate of potassium. Two or three times its volume of ether are next added, and the whole shaken up briskly for some time. After subsidence, the ethereal solution is poured off, and allowed to evaporate spontaneously, whereby a residue is left of more or less well crystallised strychnia. This may be further purified by moistening it with strong sulphuric acid, and heating it for some time in a water bath, then diluting with water, supersaturating the acid liquid with potash, again extracting with ether, and evaporating. The final product may be examined under the microscope, by the colour and physiological tests, and by the tongue. In the above process, chloroform or benzole may be substituted for the ether.

§ XII.—MORPHIA.

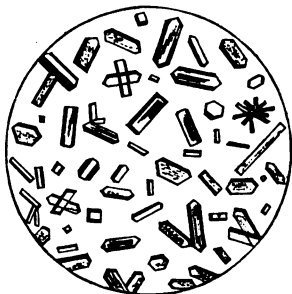
(150.) IN PURE STATE.

α. Nature, solubility, &c. Morphia is a vegetable alkaloid, having the formula $C_{17}H_{19}NO_3$. It neutralises acids to form salts, one of which, the meconate of morphia, exists largely in opium, and is the source from which the alkaloid is obtained, Morphia is readily soluble in hot, less so in cold alcohol, very sparingly soluble in ether, and almost insoluble in water, save in the presence of acids. Its ordinary salts dissolve readily in water to form solutions, which are precipitable by a great number of reagents, including most of those which precipitate strychnia. Solutions of morphia and its salts have a well marked bitter taste.

β. Appearance, &c. Morphia is usually met with in the state of acetate or hydrochlorate, which salts sometimes occur finely crystallised, but more often as imperfectly crystalline powders of a buff-tinted white colour. The alkaloid itself occurs in quadrangular prisms, frequently having two opposite edges truncated so as to produce hexagonal forms. Morphia and its salts when heated, melt, burn with a smoky flame, and leave a carbonaceous residue.

γ. Crystalline form. Morphia, when deposited by the spontaneous evaporation of its alcoholic or ethereal solution, or when slowly precipitated by caustic or carbonated alkalies from aqueous solutions of its salts, occurs in the form of variously modified prismatic crystals, as shown in fig. 52.

Fig. 52.



δ. Precipitation by potash. A single drop of potash added to the somewhat concentrated solution of a morphia salt, produces after some time, or on brisk stirring, a white precipitate

of morphia, very soluble in excess of the precipitant, but reproducible by an absorption of carbonic acid from the air.

ε. Coloursation by nitric acid. Strong colourless nitric acid added, in considerable quantity, to the cold solution of a morphia salt, produces a deep orange-red colouration. Or a drop or two of the acid may be added to a little powdered morphia or morphia salt, on a watch-glass or capsule, when an intense colouration will be at once developed. But the behaviour is not peculiar to morphia.

ζ. Coloursation by a persalt of iron. A drop or two of a carefully neutralised solution of perchloride of iron, added to a morphia solution, or to the dry alkaloid or salt, produces a deep blue colour, rendered bluish green by any excess of the iron solution.

η. Decomposition of iodic acid. A few drops of aqueous iodic acid produce, in solutions of morphia, a brown discolouration, only in part due to the liberation of iodine. Ammonia, added after a little while, deepens the colour considerably. A piece of starched paper, dipped into the coloured liquid before it has been treated with ammonia, acquires a purple colour, save when the quantity of the alkaloid is very small.

θ. Reduction of bichromate of potassium. Strong sulphuric acid produces no colouration with morphia salts or solutions. But a little bichromate of potassium solution dropped carefully on to the sulphuric acid mixture is quickly reduced with production of a bright green colour.

(151.) OPIATE LIQUIDS.

In the examination of liquids supposed to contain opium, the presence of both morphia and meconic acid is usually sought for. The last named body is not indeed poisonous, but is characteristic of opium, and possessed of well marked properties.

a. Preliminary test for morphia with nitric acid. Strong nitric acid, added in considerable quantity to an opiate liquid, will often produce a very perceptible darkening or even a distinct orange-red colouration. If necessary, the original liquid may have its colour reduced by moderate dilution with water before the addition of the acid.

β. Preliminary test for meconic acid with perchloride of iron. To the opiate solution reduced to a pale colour by dilution with even a very large proportion of water, a few drops of perchloride of iron are added, when, if meconic acid be present even in small quantity, a distinct reddening of the liquid will be produced.

γ. Precipitation of meconate of lead. The opiate liquid acidulated with acetic acid, is treated with acetate of lead, so long as a precipitate continues to be produced, when the whole is well agitated, and after partial subsidence, thrown upon a wet filter. Meconate of lead remains as an insoluble deposit on the paper, while the filtrate contains acetate of morphia, together with the excess of acetate of lead.

δ. Production of meconic acid from precipitate. The precipitate having been thoroughly washed with water, is boiled for some minutes with a small quantity of dilute sulphuric acid, and the mixture thrown upon a filter, whereby a solution of meconic acid is obtained. Or the washed precipitate suspended in a little water may be treated with excess of sulphuretted hydrogen gas, and the clear liquid filtered off, and gently evaporated.

ε. Test for meconic acid by persalt of iron. The filtrate produced by either of the above means is then tested with a few drops of perchloride of iron, which should produce a dark logwood red colour, through the formation of meconate of iron. The red liquid does not alter its colour on boiling, in which respect it differs from the similarly coloured solution of peracetate of iron, neither is it bleached by treatment with corrosive sublimate, in which respect it differs from the similarly coloured solution of sulphocyanate of iron.

ζ. Separation of morphia. Through the liquid filtered from the precipitate of meconate of lead, but having more or less acetate of lead in solution, a current of washed sulphuretted hydrogen is passed, until the smell of the gas is persistent even after agitation, when the whole is thrown upon a filter. The clear filtrate containing acetate of morphia, is evaporated down to a small bulk, supersaturated with carbonate of potassium, and agitated with an ethereal solution of acetic ether. After subsidence, the ethereal solution is poured off and allowed to evaporate spontaneously, when there is left a residue of morphia in more or less well defined crystals, to which the several tests for the alkaloid can be successfully applied.

CHAPTER IV.

ANIMAL CHEMISTRY.

§ I.—COMPOSITION OF TISSUES, &c.

(152.) ORGANIC AND MINERAL CONSTITUENTS.

a. The animal fluids and tissues consist of water together with a certain amount of solid matter. When an animal tissue or fluid is kept for some time at, or a little above, the temperature of boiling water, its aqueous portion evaporates off more or less completely, leaving the dry solids behind. This residuum, when heated upon platinum foil, undergoes combustion; some of its constituents are dissipated, and a black carbonaceous mass remains. If this carbonaceous residue be further heated for some time, especially in a current of air, the black colour will gradually disappear, and a white ash, fusible or infusible according to circumstances, will be left upon the foil. The components of dried animal matter are thus separated into two classes; one comprising the substances which are destroyed by fire, and which are called the organic constituents; the other comprising the substances which resist the action of fire, and which are called the inorganic constituents, or more simply the ashes. This distinction, however, is not absolute.

The organic components of animal matter consist principally of—

CARBON.

HYDROGEN.

OXYGEN.

NITROGEN.

SULPHUR and

PHOSPHORUS.

These are also called the elementary or *ultimate* principles of organic bodies.

The ashes consist principally of—

SODIUM.	SULPHURIC ACID.
POTASSIUM.	PHOSPHOBIC ACID.
CALCIUM.	CARBONIC ACID.
MAGNESIUM.	CHLORINE.
IRON.	FLUORINE.

Nearly all animal products are composed of both organic and inorganic constituents. Some few substances, however, pertain almost entirely to one class: thus, while the enamel of the teeth contains scarcely any organic matter, some of the crystals of uric acid met with in the urine afford scarcely any ash.

In animal tissues or fluids, the ultimate organic elements are combined with one another in a variety of ways, constituting definite compounds, which are known as *proximate* organic principles: thus in urine we may have all the above-mentioned ultimate principles united with one another, to form the proximate principles, urea, uric acid, sugar, albumen, &c.

The muscular tissue is a very suitable material to be employed for the demonstration of the principal organic and mineral constituents of animal bodies: the same general plan is adopted in other instances.

(153.) ULTIMATE ORGANIC CONSTITUENTS.

a. Desiccation. The flesh or other tissue is cut into small pieces and dried in a water bath until it ceases to lose weight. By this means it is divided into an aqueous portion which has evaporated, and a solid portion which remains. Nearly all animal matters behave in a similar way; but nitrogenous substances having an alkaline reaction, give off water containing a variable amount of ammonia.

β. Destructive distillation. A few fragments of the dried

flesh are placed in a reduction tube, into the mouth of which are inserted a narrow strip of red litmus paper, and a similar strip of lead paper. On applying the heat of a spirit-lamp, water will condense in the upper part of the tube, proving the presence of oxygen and hydrogen in the flesh—a smell of ammonia will be given off, and the litmus paper become blue, results indicating the presence of nitrogen—the lead paper will become blackened, showing the presence of sulphur—and lastly, a black mass consisting chiefly of carbon will remain in the tube.

γ. Incineration. If some of the dried flesh be heated upon a piece of platinum foil, or in a shallow capsule, it will swell up, burn with a smoky flame, and leave an abundant carbonaceous residue. On continuing the application of heat for some time, the carbon will gradually burn away. Its disappearance may be facilitated by occasionally pulverising the coherent residue resulting from the ignition. Throughout the process the temperature should not exceed, or indeed scarcely arrive at, a full red heat. As soon as a pale grey or ochry red ash is produced, the heating may be discontinued.

δ. Detection of nitrogen. A little of the finely divided dry substance is intimately mixed with ten or twelve times its bulk of soda-lime (made by slacking quicklime with caustic soda solution) and the mixture heated in a reduction tube, whereby ammonia is given off, recognisable by its smell and reaction on test-paper.

ε. Detection of sulphur. Bodies of a moderately light colour may be tested for sulphur by boiling them in aqueous potash, to which solution of acetate of lead has been added in quantity insufficient to render the liquid permanently opaque. Should the substance so treated contain sulphur it will become stained of a brown or black colour, which cannot be removed by subsequent washing with water. If a substance is stained in the above manner when boiled in a potash solution of lead, and is scarcely or not at all deepened in colour when boiled in a solution of pure potash (free from lead), the presence of sulphur is certain.

In the case of bodies readily soluble in potash water, the results are not quite so characteristic.

ζ. Deflagration with nitre. A little of the dried and finely divided animal matter is mixed with about an equal bulk of powdered nitre, and the mixture projected in small portions at a time into a porcelain crucible kept at a red heat. Deflagration immediately takes place, and, in the fused residue, the presence of carbonic, sulphuric, and phosphoric acids, resulting from the oxidation of carbon, sulphur, and phosphorus respectively, may be ascertained by the usual tests.

In performing the above experiments, very different results will be obtained with different substances, such, for instance, as pieces of flannel, hard white of egg, refined gelatine, sugar, fat, &c.

(154.) ASH OF ANIMAL MATTER.

α. A small portion of the ash, resulting from the incineration of any kind of animal matter, is placed on a watch-glass, moistened with water and examined by test-papers. Should it not have an acid reaction, a drop or so of nitric acid is to be added, and any effervescence due to carbonic acid carefully noted. The remainder of the ash is boiled in a small quantity of water for some time, a few drops of carbonate of ammonium solution added, the whole thrown upon a filter, and the filtrate set aside for examination. The residue is then well washed with water, boiled in a little hydrochloric acid with which a few drops of nitric acid have been mixed, the liquid evaporated just to dryness, diluted with water, and filtered. In this manner an aqueous and an acidulous solution are obtained, containing respectively :—

Aqueous Solution.

POTASSIUM.
SODIUM.
SULPHATES.
CHLORIDES.
PHOSPHATES.

Acid Solution.

IRON PEROXIDE.
CALCIUM.
MAGNESIUM.
PHOSPHATES.

β. Treatment of acid solution. A little of this solution may be examined for phosphoric acid by molybdate of ammonium (par. 100 *δ.*), and another portion tested for iron by ferrocyanide or sulphocyanate of potassium (par. 77). To the remainder of the solution acetate of ammonium is added, and, in the event of there being no decided reddening produced, a little perchloride of iron also. The whole is then boiled for some time, whereby a red precipitate of basic phosphate of iron is thrown down, the deposition of which may sometimes be facilitated by the careful addition of ammonia in quantity not sufficient to produce neutrality. The boiling liquid is next filtered, whereby a clear colourless solution should be obtained, perfectly free from both iron and phosphoric acid. Excess of oxalate of ammonium added to this solution throws down a precipitate of oxalate of calcium. The resulting turbid mixture, having been well shaken or stirred, is set aside for a little while, and, after partial subsidence, passed once or twice through filtering paper, when magnesium may be tested for in the clear liquid by means of ammonia and phosphate of ammonium.

γ. Treatment of aqueous solution. Separate portions of this solution, acidified with nitric acid, may be tested for sulphates by chloride or nitrate of barium (par. 95 *α.*); for chlorides, by nitrate of silver (par. 96 *α.*); and for phosphates, by molybdate of ammonium, or by sulphate of magnesium and ammonia (par. 100 *δ. α.*). Or a single portion of the acidified solution may be tested with a few drops of nitrate of barium to precipitate sulphates; then filtered and treated with excess of nitrate of silver to precipitate chlorides; then again filtered and carefully neutralised with dilute ammonia to throw down the yellow phosphate of silver which the previously free nitric acid held in solution.

The remainder of the liquid has to be evaporated to dryness, and the residue, after gentle ignition, dissolved in a small quantity of water. The solution, filtered if necessary, and acidulated with hydrochloric acid, is carefully evaporated down in a watch-

glass or capsule, when cubes of common salt will crystallise out, showing the presence of sodium. The mother liquor from these crystals is then to be treated with perchloride of platinum and alcohol, when, on stirring, a crystalline yellow precipitate of platino-chloride of potassium will be deposited. Or the solution, acidified with hydrochloric acid, may be treated at once with perchloride of platinum and alcohol, the yellow liquid filtered from the potassium precipitate and evaporated down, when yellow crystals of platino-chloride of sodium will make their appearance.

§ II.—NORMAL URINE.

(155.) GENERAL PROPERTIES.

a. Appearance, &c. Healthy human urine is an aqueous liquid in which various compounds, organic and mineral, are dissolved, and certain other substances held in suspension. It has an amber colour, a slightly acid reaction, a characteristic though not powerful odour, and a sp. gr. usually ranging within a few degrees of 1020. The substances suspended in urine are epithelium and mucus. Its dissolved organic constituents are urea, uric acid, and hippuric acid, with colouring and other extractive matters. Its inorganic or mineral constituents are sodium, potassium, calcium, and magnesium, in the form of phosphates, sulphates, and chlorides. The student is expected to identify these several substances, to make himself acquainted with their characteristic appearances, and to realise their principal reactions. A good quarter-inch object-glass is requisite for microscopic examination.

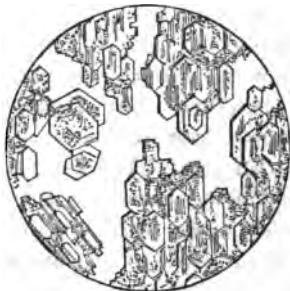
β. Mucus, epithelium, &c. Recent urine set aside for some little time in a glass vessel gradually deposits a loose flocculent sediment, readily visible upon holding the specimen between the eye and the light. When examined microscopically it is seen to consist of epithelial cells, derived from different portions of the urinary apparatus, together with granular or mucus-corpuscles. By filtration, these suspended urinary constituents

remain on the filtering paper as a scarcely visible deposit, while the urine itself passes through perfectly bright. On gently drying the filtering paper, the deposit assumes a varnish-like aspect.

(156.) UREA.

a. Its detection. A little of the filtered urine, concentrated by careful evaporation in a watch-glass, and treated with a few drops of strong colourless nitric acid, yields either at once or very speedily a crystalline deposit of nitrate of urea. This deposit, when examined under the microscope, is seen to consist of delicate six-sided plates, superimposed upon one another so as usually to prevent more than three or four of the sides of any one crystal being recognisable, as shown in fig. 53. The nitrate of urea may be formed in a watch-glass and then transferred to a slide, or it may be dissolved in water and recrystallised on a slide, or nitric acid may be added to some concentrated urine previously placed in the field of the microscope, and the actual process of crystallisation observed.

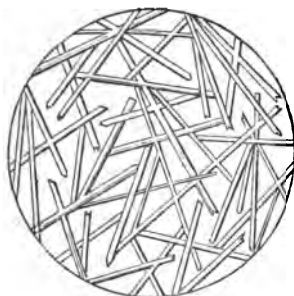
Fig. 53.



β. Preparation of urea. A couple of ounces or so of fresh and filtered urine are evaporated on a water-bath or gently heated sand-bath to a syrupy consistency, and a quantity of strong colourless nitric acid about equal in bulk to the concentrated urine added thereto, when the mixture, upon cooling, becomes semi-solid from the formation of nitrate of urea. The crystalline mass is drained on a tile, or pressed between several folds of blotting paper, then dissolved in a little warm water, and the resulting solution treated with an excess of carbonate of barium. Upon concentrating the filtered liquid, nitrate of barium crystallises out first, while urea remains in the mother liquor, which is evaporated to dryness over a water-bath. From

this residue warm alcohol extracts the urea, and, on cooling or slow evaporation, deposits it in the form of long flattened prismatic crystals, as shown in fig. 54.

Fig. 54.



γ. Properties of urea. Urea behaves in some respects like an organic base, being capable of uniting with certain acids, notably the nitric and oxalic acids, to form salts. It dissolves readily in water and alcohol, producing solutions which are neutral to test-paper. The formula of urea is $\text{CH}_4\text{N}_2\text{O}$, and that of nitrate of urea $\text{CH}_4\text{N}_2\text{O}.\text{HNO}_3$. Urea is isomeric with cyanate of

ammonium NH_4CNO , which undergoes spontaneous conversion into it; and also isomeric, if not identical, with carbamide $\text{N}_2\text{H}_4(\text{CO})$. Heated with water under pressure, it is transformed into carbonate of ammonium, thus: $\text{CH}_4\text{N}_2\text{O} + 2\text{H}_2\text{O} = (\text{NH}_4)_2\text{CO}_3$. The same change takes place spontaneously in putrefying urine, and is also brought about by acting on urea with strong potash or sulphuric acid, except that the resulting carbonate of ammonium is then broken up by the reagent employed. 1,000 parts of urine contain, on the average, about 15 of urea.

(157.) URIC ACID.

a. Its detection. This compound occurs but in small quantity in healthy human urine, 100 parts of which contain, on the average, not more than half a part of uric acid. In order to detect its presence, a couple of ounces or so of filtered urine are reduced to one-half the original bulk by evaporation, a little hydrochloric acid added to the concentrated liquid, and the whole set aside in a cool place for some hours, when the interior of the vessel will be found studded with small brown crystals of impure uric acid. In

the case of urine having a moderately high specific gravity, concentration is unnecessary. After pouring off the supernatant liquid, the crystals are detached, washed with water, and dissolved in a few drops of warm potash. The resulting solution of urate of potassium is then filtered, and acidulated with hydrochloric acid, whereby a crystalline precipitate of uric acid is thrown down, which may be examined microscopically and by the action of nitric acid, as described below.

β . Preparation of uric acid. Uric acid cannot well be prepared, in any quantity, from normal human urine. But it may be easily obtained from the common brickdust urinary deposit, collected on a filter and washed with water; or from powdered uric calculi; or the excreta of serpents. Any one of these substances is boiled with caustic potash, the solution diluted with water, filtered, and supersaturated with hydrochloric acid, whereby a very considerable opacity is at first produced, which, however, speedily disappears, and is replaced by a dense crystalline precipitate, from which the supernatant liquid may be readily poured off.

γ . Properties of uric acid. Uric acid furnishes two classes of salts, acid and neutral, and is consequently dibasic. Its formula is $C_5H_4N_4O_3$. The formula for a scarcely soluble or acid urate is $C_5MH_3N_4O_2$, and that for a soluble or neutral urate $C_5M_2H_2N_4O_3$. Uric acid itself is extremely insoluble, both in water and alcohol; but is soluble in alkaline solutions, forming neutral urates, and reprecipitated therefrom on the addition of an acid. It always occurs in the crystalline state, the appearance of the crystals, however, being very various. Occasionally the acid is met with in its normal form of the rhombic prism, more frequently in rhombic plates with the obtuse angles more or less rounded off, or in acuminate doubly-convex lozenge-shaped plates, or in elongated flat plates with excavated ends. Some of the forms of uric acid crystals are shown in figs. 55 and 56. Uric acid dissolves readily, with effervescence, in nitric acid, and

on evaporating the solution to dryness an amorphous pinkish residue is left. This, when moistened with ammonia, assumes a

Fig. 55.

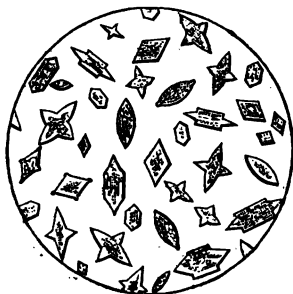
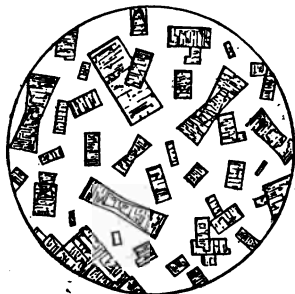


Fig. 56.



fine crimson colour, which is changed to violet on the addition of a small quantity of caustic potash.

(158.) HIPPURIC ACID.

a. Its preparation. 'Although this acid exists, in healthy human urine, in nearly the same proportion as uric acid, yet its presence therein does not so readily admit of demonstration. It can, however, be easily procured from the urine of herbivora, and from that of patients who have been taking benzoic acid as a medicine. To prepare it from either of these sources, the recent filtered urine is evaporated down to one-fourth of its bulk, and then treated with an equal volume of ordinary hydrochloric acid, when, on cooling, long prismatic needles of impure hippuric acid crystallise out. These, after being washed with a little cold water, are dissolved in boiling water, and the solution set aside to crystallise.

β. Properties of hippuric acid. The hippuric is a monobasic acid, represented by the formula $C_9H_7NO_3$. It is soluble in water, alcohol, and ether. The crystals obtained by cooling the hot aqueous solution on a slide consist of delicate

prisms, often presenting the appearance of elongated six-sided plates, as shown in fig. 57.

By prolonged boiling with concentrated hydrochloric acid, hippuric or glyco-benzoic acid absorbs a molecule of water, and breaks up into benzoic acid and glycocine or sugar of gelatine, thus: $C_9H_7NO_3 + H_2O = C_7H_5NO_2 + C_2H_5NO_2$. The larger proportion of the former product is dissipated by evaporation; but the glycocine may be detected

Fig. 57.



by adding to the liquid a drop or so of aqueous sulphate of copper and an excess of potash, whereby a deep blue-coloured solution is produced, unaffected by ebullition.

(159.) COLOURING AND EXTRACTIVE MATTERS.

a. Purpurine, &c. When healthy urine is boiled in a test-tube with about one-fourth its bulk of hydrochloric acid, a deep brownish-purple colour is produced, due to the metamorphosis of a peculiar highly carbonised pink colouring matter, known as purpurine. The common pink deposits of alkaline urates owe their colour to this purpurine, which has a great tendency to become precipitated with them. Hence, when perfectly white urate of ammonia, *boa's* excrement, for instance, is boiled in urine containing much purpurine, it is deposited on cooling of a pink colour, from its carrying down some purpurine with it. When these coloured deposits, natural or artificial, are boiled in alcohol, the purpurine is dissolved, forming a pinkish-red solution. The relation in which purpurine stands to the yellow colouring matter of urine is not satisfactorily established.

β. Extractives. The remaining organic urinary constituents are called by this name. They generally amount to about one per cent. of the urine. Included among them is some principle containing sulphur in an unoxidised form, also kreatine and

kreatinine, substances derived from the oxidation of muscle, and found more largely in the juice of flesh. Schunck has noticed in urine the habitual presence of a substance, allied to indican, decomposable by an absorption of water into grape sugar and indigo-blue.

(160.) MINERAL SALTS.

a. Ash of urine. This may be examined according to the directions previously given for the analysis of animal ashes (par. 144). It will be usually found to contain calcium, magnesium, sodium, and a small quantity of potassium, in the form of carbonates, sulphates, phosphates, and chlorides. The residue left after the ash has been acted upon by acid, consists principally of carbon. When this residue is ignited for some time, a minute white ash remains, which is said to contain silica and fluorine.

β. Most of the inorganic constituents of urine may be detected in the secretion itself after simple filtration. On the addition of ammonia a white precipitate is produced, which, when examined microscopically, is seen to consist of amorphous granules of phosphate of calcium and characteristic stellate feathery crystals of ammonio-phosphate of magnesium. If healthy urine, especially that passed after a

Fig. 58.



night's rest, be evaporated very gradually upon a glass slide, minute octahedral crystals of chloride of sodium may be detected by microscopic examination. Generally speaking, however, the common salt is seen in the form of very complex right-angled crosslets (stauroid crystals), arranged somewhat like *chevaux de frise*. On the axes and extremities of these forms, octahedrons may occasionally be recognised by a good defining glass. Fig. 58 shows one of the many appearances presented by carefully evaporated urine.

Sulphuric and hydrochloric acids may be detected in urine by the usual tests.

§ III. ABNORMAL URINE.

The abnormal constituents of urine which do not necessarily give rise to deposits, are albumen, sugar, biliary matters, and fat.

(161.) ALBUMINOUS URINE.

a. Appearance, &c. The general appearances of albuminous urine vary considerably. Sometimes it presents nothing unusual in its aspect; often the flocculent deposit formed by repose is larger in amount than that of the healthy secretion; occasionally the urine has a faintly opalescent appearance, not removable by filtration; and very frequently it is met with, black, brown, or red, from the presence of altered blood, with or without the occurrence of a deposit of blood globules. Albuminous urine when shaken retains the froth for a long time; its specific gravity is very variable.

β. Test by boiling. Some of the suspected urine is boiled in a test-tube, when should albumen be present, a turbidity will be produced, the amount of which may vary from a faint cloud to a bulky precipitate rendering the urine nearly solid. *Impediments.*—a. Albumen when dissolved in alkaline fluids is not necessarily deposited upon boiling, the formation or non-formation of a precipitate having reference to the relative quantities of albumen and alkali respectively present. Therefore, in testing an alkaline urine for albumen, the liquid should first be rendered very faintly acid with acetic acid. b. A previously opaque condition of the urine interferes with the action of this test. This interference may be lessened, if not removed, by filtering the urine before applying heat thereto. In urine containing deposits, the clear liquid can generally be poured off and tested separately. But deposits of urates do not impede the action of this test, as, upon heating the urine, their dissolution takes place before the precipitation of any albumen. c. Albuminous urine, to

✓ which a very minute quantity of nitric acid has been added, is not rendered turbid by heat. *Fallacy*.—Upon boiling certain varieties of urine, a precipitate of the earthy phosphates occasionally takes place, which can, however, be distinguished from that of albumen by the addition of a little dilute nitric acid, whereby the former is dissolved, the latter unaffected.

✓ γ. Nitric acid test. On adding nitric acid to albuminous urine, a white turbidity is produced, varying in amount with the proportion of albumen present. *Impediments*.—*a*. The reagent should be added drop by drop, since a minute quantity does not cause any precipitate, and a great excess dissolves any precipitate which may have formed. *b*. When the urine is opaque, it should be rendered slightly alkaline with potash, agitated, filtered, and then tested with nitric acid. *Fallacies*.—*a*. In some varieties of urine the addition of nitric acid produces a precipitate of uric acid, which, however, speedily shrinks very much in bulk, and, when examined microscopically, is seen to be crystalline, whereas the deposit of albumen is amorphous. *b*. Nitric acid produces a whitish turbidity in the urine of patients who have been taking copaiva, cubebs, and probably other oleo- and resinous medicines. But while the precipitate of albumen subsides to the bottom of the test-tube in the course of a few hours, the precipitate of oleo-resinous matters remains suspended in the urine for two or three days. Moreover, inquiry can always be made on these points.

δ. Ferrocyanide of potassium test. Solution of ferrocyanide of potassium added to albuminous urine previously acidulated with acetic acid, throws down a white precipitate. *Impediment*.—The mere addition of acetic acid to urine occasionally produces a precipitation of mucus, in which case the acidulated urine must be filtered before being tested with the ferrocyanide.

!(162.) SACCHARINE URINE.

✓ α. Appearance, &c. Saccharine urine cannot be distinguished by the eye from the normal secretion. It has generally a high specific gravity, a rather fragrant odour, and when agitated

retains its froth for some time. It is said to have a distinctly sweet taste. Minute traces of sugar exist habitually in normal urine.

β. Potash test. To the suspected urine, an equal bulk of the ordinary solution of caustic potash is added, and the whole boiled; whereon a deep orange-brown, frequently almost black, colour is produced if sugar be present in abnormal quantity.

Fallacies.—Should a deep brown colour be produced, the test is free from fallacy; but there are many specimens of non-saccharine urine which, when boiled with caustic potash, acquire a tolerably dark sherry colour. Moreover, caustic potash frequently contains lead, and this impure reagent, acting upon the sulphur of ordinary urine, or more decidedly in albuminous specimens, will occasion a brown discoloration. The potash should, therefore, be first tested for lead. ✓

γ. Copper test. The urine is mixed with about half its bulk of caustic potash solution, whereby a precipitate of earthy phosphates is produced, which, in questionable cases, should be separated by filtration. To the alkaline liquid, filtered or unfiltered, a drop or two of a dilute solution of sulphate of copper is added, when, if sugar be present, the greenish-blue precipitate first thrown down will disappear upon agitation, forming a deep blue-coloured liquid. On heating this liquid, and even before it arrives at the boiling point, a red or orange-coloured precipitate will be formed, characteristic of sugar. *Fallacies.*—Although the precipitate produced by the addition of sulphate of copper to a mixture of normal urine and caustic potash does not disappear upon agitation, still the disappearance of the precipitate with formation of a blue liquid is no proof of the presence of sugar. Moreover, the application of heat should not be continued after the blue liquid has acquired a boiling temperature, as many substances by prolonged ebullition effect a deposition of the red suboxide of copper. ✓

δ. Modified copper test. Instead of taking separate solutions of sulphate of copper and potash, a ready-made alkaline solution of tartrate of copper may be added to the suspected

urine, and the whole heated to the boiling point, when the production of an orange precipitate of suboxide of copper will show the presence of sugar. The test solution is made by dissolving about 20 grains of sulphate of copper and 40 grains of neutral tartrate of potassium in an ounce of the officinal solution of potash, whereby a clear deep blue liquid should be produced, which may be filtered if necessary.

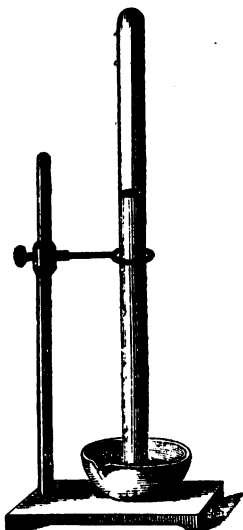
ε. Tin test. The reductions of salts of tin, bismuth, silver, and chromic acid by grape sugar at a moderate heat, have been made the bases of separate tests. The tin test is best performed by having ready-prepared strips of merino or other woollen (not cotton or linen) tissue dipped in solution of dichloride of tin and then dried in a water-bath. On moistening one of these strips

with diabetic urine, and holding it near the fire, or otherwise heating it to a temperature of about 300° F., a brownish-black coloration quickly makes its appearance. This is a convenient clinical test, and one of great delicacy.

ζ. Fermentation test. Ordinary yeast, or the dried German yeast, is mixed with water, and a long test-tube completely filled with the suspected urine, to which a little of the yeast liquid has been added. The tube is then closed with the thumb, and inverted in a saucer containing a little of the urine under examination, so that no air may enter the tube; and the whole set aside in a tolerably warm situation. The temperature ought not to be below 70° Fahrenheit. Should

the urine be saccharine, minute air-bubbles will speedily make their appearance, and in the course of an hour or so a very definite quantity of gas will occupy the upper part of the tube,

Fig. 59.



as shown in fig. 59; but other forms of apparatus may be employed. The sporules and thallus of the sugar fungus, or yeast plant, are said to be recognisable in stale saccharine urine.

(163.) BILIARY URINE.

a. Appearance, &c. Biliary urine has a yellowish-brown colour, and a persistent bitter taste. It is doubtful whether the urine as voided ever contains more than a trace of the true biliary salts, for the detection of which substances only is Pettenkofer's test adapted (*vide par.* 178); but Heller's test, and the nitric acid test, react upon the colouring matter of bile, which not unfrequently finds its way into the urine.

Fig. 60.



β. Nitric acid test. A little of the urine, previously concentrated if necessary, is poured on to a white plate, so as to form a thin layer, upon which a few drops of strong nitric acid are then let fall. Where the acid comes into contact with the biliary urine, a peculiar play of colours is produced—green, pink, violet, and yellow, being readily recognisable. Or a mixture of the urine with dilute nitric acid may be carefully poured on to some strong sulphuric acid, when the characteristic colouration will take place at the junction of the two liquids, as seen in fig. 60.

γ. Heller's test. For the application of this test the urine is required to contain albumen; hence it must be mixed, if necessary, with a little diluted white of egg, serum of blood, or some other urine containing albumen. Nitric acid is then added, when, if bile be present, the precipitate will have a faintly bluish or greenish colour; but the test is not very satisfactory.

(164.) FATTY URINE.

a. Fat globules, &c. The conditions in which fat occurs in the urine have not been ascertained with any degree of precision.

Occasionally, when examining the ordinary flocculent deposit of urine by means of the microscope, isolated fat globules may be recognised. Fat occurring only in this state is believed by some observers to be necessarily of extraneous origin. In some forms of Bright's disease the fibrinous casts of tubes and the epithelial cells, particularly those derived from the kidney, are seen loaded with fat, and at the same time isolated fat globules may be detected.

β. Chylous urine. Occasionally this curious variety of urine is met with: the secretion is more or less opaque, always contains albumen, frequently gelatinises on cooling, and, when examined microscopically, displays an abundance of minutely divided granular matter, and a few granular cells similar to those found in the chyle, but no fat globules. Fat, however, may be readily obtained by agitating the urine with ether, and evaporating the ethereal solution.

γ. Kiestein-urine. In the urine of pregnant women the so-called kiestein may be recognised. The secretion has generally an acid reaction, and by repose becomes faintly opaque. In the course of two or three days a fat-like scum rises to the surface, remains there for two or three days, and then sinks to the bottom of the vessel, the urine becoming at the same time ammoniacal. When this scum is examined microscopically, it is seen to consist of crystals of triple phosphate, with a few fat globules, imbedded in a dense granular matter, which appears to be of an albuminous character, containing, however, minutely divided fat. Kiestein urine, by keeping, frequently evolves a powerful odour of putrescent cheese. Its occurrence is no longer regarded as positively diagnostic of pregnancy.

§ IV. URINARY DEPOSITS.

These may be distinguished into chemical compounds either crystalline or amorphous, and structural or organised compounds.

(165.) CHEMICAL DEPOSITS.

a. Urates or lithates.—*Every deposit which disappears upon the application of heat, consists of uric acid in combination with various bases.* These urates are sometimes white, but generally more or less coloured with purpurine, which may be partially extracted from them by means of boiling alcohol. They form a bulky deposit, nearly always amorphous, occasionally appearing in the form of minute spheres with protruding spicula. Urate sediments dissolve readily on the addition of caustic potash, the solution when boiled giving off ammonia. Moreover, they yield a residue of murexide, when treated with strong nitric acid, evaporated to dryness, and held over the vapour of ammonia. The urine in which these deposits occur has generally an acid reaction.

β. Earthy phosphates.—*Nearly every deposit which disappears upon the addition of hydrochloric acid consists of earthy phosphates.* These salts form a bulky opaque white deposit, which, unless associated with mucus, is easily diffusible upon agitation. The urine itself is mostly alkaline, or neutral, or but very faintly acid. Deposited phosphate of calcium is usually amorphous, save in acid urine, where it sometimes occurs in peculiar radiated or penniform crystals. The ammonio-phosphate of magnesium crystallises in variously modified triangular prisms, often simulating irregular six-sided plates, as shown in fig. 61; and in very ammoniacal urine is met with in the form of stellate feathery crystals.

Fig. 61.



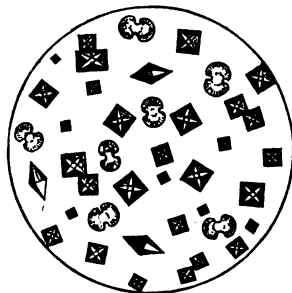
In some cases this deposit is seen as an iridescent pellicle, and occasionally it remains for a long time suspended in the urine.

γ. Uric acid.—*Every obviously crystalline deposit, having a*

distinctly yellow or red colour, consists of uric acid. The colour, specific gravity, and acidity of urine yielding uric acid sediments are generally rather above than below the average. The deposit is compact in its appearance, and subsides quickly after agitation. It is readily soluble in potash, from which solution it is reprecipitated on the addition of hydrochloric acid. When treated with strong nitric acid, evaporated to dryness, and held over the vapour of ammonia, it yields murexide. Uric acid deposits are very rarely indeed devoid of colour. They are met with in variously modified crystalline forms (*vide par.* 147).

δ. Oxalate of calcium. This salt seldom if ever forms a distinct sediment. It may be detected by allowing the urine to stand at rest for some time, and then pouring away all but the

Fig. 62.



last portions, which must be examined microscopically. It occurs in well-marked octahedrons, the crystals generally appearing to have a square outline, and their opposite angles being connected by markings, as shown in fig. 62. The sediment may be rendered apparent to the naked eye, by warming the residue of urine, left after pouring away the greater portion, giving it a rotatory motion, and allowing it to stand for

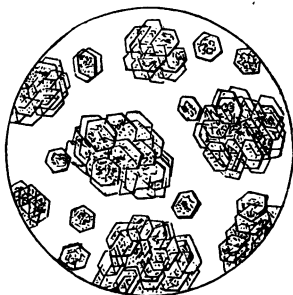
a few minutes. Then on pouring off the remainder of the urine, and replacing it by water, a white glistening deposit becomes visible. The deposit of oxalate of calcium is scarcely at all affected by cold potash. It dissolves without effervescence in dilute hydrochloric acid, forming a solution which effervesces on the addition of a little peroxide of manganese. There are occasionally found associated with octahedrons of oxalate of calcium, certain dumb-bell-shaped crystals, said to consist of oxalurate of calcium. They are insoluble in potash, soluble in hot hydrochloric acid, and when incinerated leave a residue of carbonate of

calcium. Deposits of uric acid and phosphates have been occasionally met with in the dumb-bell form.

e. Cystine. This deposit occurs somewhat rarely, in the form of a bulky, easily diffusible deposit, resembling in its appearance the white or pale lithates. When examined microscopically it is seen to consist of rosette-like plates, in which a hexagonal outline may sometimes be recognised. After pouring off the supernatant urine, the sediment will be found to be insoluble in acetic acid, soluble in hydrochloric acid, and very soluble in ammonia. When the ammoniacal solution is allowed to evaporate

spontaneously on a slide, very well-defined, transparent hexagonal plates crystallise out, as shown in fig. 63. Cystine is remarkable for containing twenty-six per cent. of sulphur, so that when cystic urine is boiled with a solution of acetate of lead to which potash has been added in sufficient excess to dissolve the precipitate at first thrown down, the whole becomes nearly black, from the formation of sulphide of lead. Cystic urine has when recent an aromatic, when decomposing a very foetid odour. The formula of cystine is $C_3H_7NSO_3$.

FIG. 63.



(166.) ORGANISED DEPOSITS.

a. Normal sediment. A flocculent deposit, varying slightly in quantity and appearance, always separates from urine by repose. When examined microscopically it may show, in addition to the usual mucus-corpuscles and epithelium-cells, torulæ characteristic of saccharine urine, casts of uriniferous tubes indicative of Bright's disease, fat globules, whether free or contained in a cell wall, vibriones, spermatozoa, blood globules, exudation globules, minute crystals of oxalate of calcium, and occasionally of uric acid, &c. &c.

β. Pus. This substance presents a different appearance accordingly as it occurs in acid or alkaline urine. In acid urine, purulent deposits sink to the bottom of the vessel, and form a greenish-yellow opaque layer, having a creamy consistency, an easy diffusibility on agitation, and a slow subsidency on repose, gelatinising when agitated with an equal bulk of caustic potash, and in fact presenting all the ordinary qualities, physical and microscopical, of pus. In alkaline urine the deposit is viscid, tenacious, ropy, not diffusible on agitation, and is mixed up with the earthy phosphates, which may, however, be separated by the action of dilute hydrochloric acid. Purulent urine is necessarily albuminous. When a deposit of pus has been

agitated with ether, the resulting ethereal solution, upon being poured off and evaporated, leaves a considerable residue of oily globules.

The pus corpuscles, which, however, can scarcely be said to exist after the action of an alkali, consist of circular granulated cells, somewhat larger than blood globules, as shown in fig. 64. When acted on by acetic acid, they swell up very considerably, their margins become

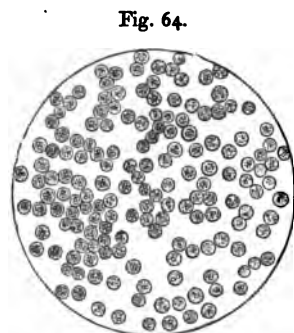


Fig. 64.

distinct, and two, three, or four small granular masses appear in their centres.

γ. Mucus. A little mucus is always met with, even in normal urine. It may be much increased in quantity without being appreciably altered in character; but oftener it occurs in the form of gelatinous masses, which sink to the bottom of the vessel, or, from the entanglement of air-bubbles, remain for a long time suspended in the urine.

Mucus deposits have a more or less marked alkaline reaction, even when the supernatant urine is acid: they do not diffuse readily by agitation, and are frequently associated with a very

considerable sediment of the earthy phosphates. The presence of even a large amount of mucus does not of itself render the urine albuminous. The microscopical characters of the mucus corpuscle are very similar to those of the pus corpuscle, but the granular character is perhaps not quite so well marked.

δ. Blood. Urine containing blood is necessarily albuminous. On allowing the urine to stand, and examining the sediment microscopically, blood globules may be recognised by their uniform size, non-granular surface, and yellow colour. The appearance of urine containing blood is very variable: its colour may be light-red, dark-red, reddish-brown, smoke-brown or scarcely altered. The sediment also varies much in its appearance: sometimes its nature is evident to the unassisted eye, at other times it occurs but in very minute quantities, and can only be identified microscopically. Urine may contain altered colouring matter of blood, and yet the blood-corpuscles escape recognition.

§ V.—CLINICAL EXAMINATION OF URINE.

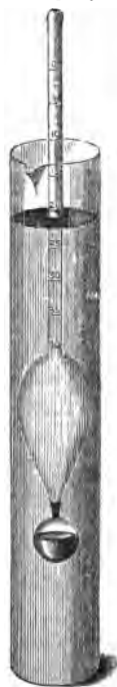
(167.) GENERAL EXAMINATION.

α. Appearance, &c. It is advisable to notice the colour, whether pale from a dilute urine, or dark from a concentrated urine, or reddish yellow from purpurine, or brownish yellow from bile, or red or brown from blood—the taste, whether sweet from sugar, or bitter from bile—the smell, whether fragrant from cystine or from sugar, or fœtid from alkalinity, with or without mucus, or from cystine—any opalescence or milkiness due to fat, kiestein, mucus, or a modification of albumen, &c.

β. Specific gravity. The specific gravity should be taken by means of a gravimeter (fig. 65.) It may be too low, from an accidentally diluted urine, or from *Diabetes insipidus*, or from certain forms of *Morbus Brightii*, &c.; too high, from a concen-

trated urine or from the presence of sugar, &c. Dr. Golding Bird pointed out the very curious coincidence, that the last two

Fig. 65.



figures, expressing the specific gravity, represent very nearly the number of grains of solid matter contained in an ounce of the secretion: thus in urine of the specific gravity 1017, every fluid ounce contains about 17 grains of solid matter. This, however, must be regarded as but a very rough approximation to the truth, trifling variations in the amount of the saline constituents of urine effecting greater alterations of density than considerable variations in its organic constituents.

γ. Quantity. In order to determine the quantity of urine passed in twenty-four hours, the patient should be caused to make water at some definite hour in the day,—say 10 A.M., the amount then voided being neglected. After this he should save all the urine he passes until 10 o'clock on the next day, at which time he must again empty his bladder, and add the contents to the specimen to be measured. He should moreover be desired always to micturate before going to the closet. By multiplying the number of ounces passed, by the last two figures of the specific gravity, an approximation will be arrived at as to the total amount of solids excreted by the kidneys in twenty-four hours.

(168.) CHEMICAL EXAMINATION.

α. Reaction to test-paper. Normal urine has a slightly acid reaction. If alkaline, it will restore the blue colour of reddened litmus-paper. Should the blue colour remain after the paper has become dry, the alkalinity is due to the presence of a

fixed alkaline salt; but should the red colour reappear on drying the paper, the alkalinity is due to ammonia.

β. Testing the urine. The supernatant urine is to be poured away from any deposit which may have formed, and be examined for albumen, sugar, purpurine, and if necessary fat and bile. Should the urine have a high specific gravity and be free from sugar, it may be examined for an excess of urea by pouring a little of the secretion into a watch-glass, and adding about two-thirds its bulk of cold concentrated nitric acid. The formation of a crystalline deposit of nitrate of urea is, with certain restrictions, indicative of an excess of that base. It is, perhaps, generally advisable to concentrate the urine slightly, before adding the acid.

γ. Testing the deposit. The appearance of the deposit generally indicates the order in which the tests, both microscopical and chemical, should be applied. The lithates are dissolved by heat, dissolved by potash, undissolved by hydrochloric acid; the phosphates are undissolved by heat, undissolved by potash, dissolved by hydrochloric acid; uric acid is undissolved by heat, dissolved by potash, undissolved by hydrochloric acid; cystine is undissolved by heat, dissolved by ammonia or potash, and dissolved by hydrochloric acid; oxalate of calcium is undissolved by heat, undissolved by potash, dissolved by hydrochloric acid. In mixed deposits the different ingredients are readily recognised by their different microscopical appearances, and by their different behaviour with the above reagents.

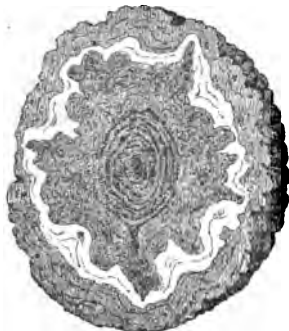
§ VI.—URINARY CALCULI.

(169.) GENERAL CHARACTERS.

a. Construction, &c. Urinary calculi are for the most part built up of concentric layers. This structural arrangement is readily seen on making a section of a calculus through its

centre. All the layers of a calculus may have the same composition, or may differ very much from one another in this respect. One single uniform layer of a calculus may be, and generally is, composed of several ingredients. The internal arrangement of a

Fig. 66.



mixed mulberry and fusible calculus, belonging to the museum of St. Bartholomew's Hospital, is shown in fig. 66 (No. 79 in catalogue).

It is probable, that if a very exact analysis were made, each of the layers of nearly every calculus would be found to contain uric acid, alkaline urates, phosphate of calcium, and ammonio-phosphate of magnesium, with or without the other constituents of calculi. Moreover, most calculi contain traces of all

the salts naturally existing in the urine, as well as of colouring matter, mucus, &c.

From these considerations it is obvious that the chemical examination of a calculus need have reference only to its general composition, and not to its exact analysis. It is important, however, to bear in mind, that even a homogeneous layer of a calculus rarely ever consists of one constituent only.

β. Appearance, &c. The general appearances, &c., of the different varieties of calculi are as follows:—*a.* Uric calculi consist of uric acid, with or without variable proportions of the alkaline urates. They have usually an uniform outline, a compact laminated structure, and an orange or yellow colour: sometimes the laminated appearance is wanting, and sometimes they have a light fawn colour, resembling the paler varieties of oxalate of lime concretions. A greater or less amount of uric acid is found in the centres of most calculi.

b. Earthy phosphates consist of phosphates of calcium and ammonio-phosphate of magnesium. It rarely if ever happens that

where one of these constituents is present the other is wholly absent. When the two exist in about equal quantities, the concretion is known by the name of the *fusible calculus*, in consequence of the readiness with which it fuses in the blowpipe flame. But when either constituent is present in great excess, this fusion cannot be effected. The distinctly laminated character appears to be more frequently wanting in the fusible than in most other varieties of calculi. Phosphatic calculi differ much in their appearance; they have usually a smooth uniform surface, and a pale, white, or even chalk-like aspect. Sometimes they are compact and hard, at other times light and friable; sometimes the layers adhere very closely to one another, and at other times are just as easily separable. The earthy phosphates may constitute the greater part of a calculus, or may be disseminated through the other constituents, or may form distinct layers: they give a more or less thick external coating to most calculi.

c. Oxalate of calcium calculi are generally recognised by their dark colour, hard compact laminated structure, and irregular surface. The term mulberry calculus does not give any idea of the degree of this irregularity. But some small oxalate of calcium calculi, known as hemp-seed concretions, have a smooth contour. Occasionally, oxalate of calcium concretions, especially when forming layers in other calculi, or when mixed with uric or phosphatic deposits, have a pale colour, and very finely laminated structure. The central portions of oxalate of calcium calculi generally contain more or less uric acid.

d. Cystine calculi are of comparatively rare occurrence. They have an irregular or oval shape, a rough and crystalline-looking surface, a fawn-brown colour when recent, and a sea-green colour when long kept. Cystine rarely ever enters into the constitution of composite calculi.

γ. Action of heat. Some of the ingredients of calculi are destructible by heat, some indestructible, as seen in the following table:—

Destructible.	{	URIC ACID.
		URATE OF AMMONIUM.
		CYSTINE.
		OXALIC ACID, from oxalate of calcium.
		AMMONIA, from triple phosphate.
Indestructible.	{	URIC ACID, from urates of calcium and sodium.
		PHOSPHATE OF CALCIUM.
		CARBONATE OF CALCIUM.
		PHOSPHATE OF MAGNESIUM, from triple phosphate.
		CARBONATE OF SODIUM, from urate of sodium.
		CARBONATE OF CALCIUM, from oxalate and urate of calcium.

If the heat be sufficiently prolonged and intense, the carbonate of calcium will become converted into caustic lime. Carbonate of calcium is not an unfrequent constituent of calculi which have undergone partial decomposition in the bladder.

(170.) PRELIMINARY EXAMINATION.

a. Pulverisation, &c. The calculus to be examined should be sawn through its centre, so as to expose its internal arrangement. Should it consist of layers obviously differing from one another each of them must be separately examined. For this purpose, a sufficient quantity of each layer may be consecutively removed by means of a pocket-knife. The determination of the nature of any one layer should be ascertained before removing a specimen of the next one. In friable calculi, great care must be exercised in obtaining specimens of the different layers. The smooth appearance of the flat surface can be readily restored by grinding, so that a calculus may be analysed without any necessary disfigurement. The specimen removed from each layer is to be reduced to a fine powder.

β. Ignition. A little of the powder is to be heated upon platinum foil, and careful attention paid to any of the following results, &c.

Charring. All urinary calculi undergo a slight amount of charring. In oxalate of calcium calculi this is very slight, and speedily disappears, leaving a bulky white pulverulent residue. In phosphatic calculi the charring is more complete, and the carbon not so easily burnt off.

Decrepitation. This is always very slight: when occurring simultaneously with the formation of a white smoke and a great degree of mobility in the heated powder, it is indicative of urate of ammonium.

Odour. Oxalate of calcium calculi do not evolve much odour when heated: most others do. The odour produced by the ignition of cystine is well marked and characteristic.

Volatilisation. Should the calculus powder burn away almost entirely, it will suffice to test for uric acid, urate of ammonium, and cystine.

Fusion. The heat of a spirit-lamp is sometimes sufficient to fuse the mixed earthy phosphates.

Alkalinity. When only the heat of a lamp has been employed, any alkalinity to test paper, shown by the moistened residue, is probably due to carbonate of sodium, derived from the ignition of urate of sodium.

Effervescence. The moistened residue is treated with a drop or two of nitric acid. Effervescence denotes the presence of a carbonate, whether originally existing as such, or derived from the ignition of oxalate of calcium or of the fixed alkaline urates; in which latter case the amount of effervescence is usually very small.

Blowpipe. Should the ordinary flame have proved incapable of fusing the ash, the acidified residue may now be dried, and strongly heated before the blowpipe; when, if the mixed earthy phosphates are present, a more or less complete fusion, or at any rate cohesion of the particles, will take place.

(171.) SPECIAL TESTS.

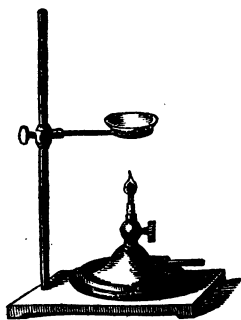
In addition to noticing the above described effects of heat, it

will often suffice to make special tests for the phosphoric, oxalic, and uric acids, without subjecting the calculus to a more complete analysis.

a. Phosphoric acid. The residue upon the foil after its ignition before the blowpipe is treated with a drop or two of nitric acid and a little water, whereby its solution is usually effected without appreciable effervescence, any carbonate of calcium having been converted into caustic lime. A drop or two of nitrate of silver solution is then added, and the mixed liquid carefully neutralised with dilute ammonia, when the production of a yellow precipitate will indicate the presence of phosphoric acid. Under certain circumstances, especially when triple phosphate constitutes the great mass of the calculus, or when the silver salt has been added in very small quantity, the precipitate is white. Or the nitric acid solution of the ignited calculus may be tested with molybdate of ammonium (*vide par.* 92 δ). Triple

phosphate are distinguished from bone earth calculi by their solubility in acetic acid.

Fig. 67.



β . Oxalic acid. Some of the original calculus powder is mixed with a drop or so of dilute sulphuric acid, and a little finely divided peroxide of manganese added, whereby any oxalic acid is at once oxidised into carbonic acid, which is liberated with effervescence.

γ . Uric acid. A little of the original calculus powder, placed in a watch-glass, is treated with a drop or two of strong nitric acid in which, if uric acid or a urate be present, it will dissolve with effervescence. On carefully evaporating to complete dryness over a small flame, as shown in fig. 67, a pinkish residue is left, which, when cold, is to be moistened with a drop of ammonia, whereby murexide will be produced with its characteristic crimson colour, convertible into violet on the addition

of a little caustic potash. Or the watch-glass may be held in the fingers as a precaution against using too strong a heat.

(172.) SYSTEMATIC ANALYSIS.

a. Solution. Some of the powdered calculus is boiled for a few minutes with a little distilled water in a test tube, the mixture thrown upon a filter, the filtrate collected apart, and the residue thoroughly washed with boiling water. The first portion of the washings may be reserved for use on an emergency. The filtrate *A* may contain urate of ammonium, urate of sodium, and urate of calcium.

The washed residue is next boiled in dilute hydrochloric acid, observation being made as to whether or not any effervescence indicative of the presence of carbonate of calcium takes place. The acid liquid is thrown on a filter, the resulting filtrate *B*, collected apart, and the residue therefrom, if any, washed with water. The acid solution *B* may contain chloride of calcium from the decomposition of the carbonate, oxalate of calcium, cystine, phosphate of calcium, and ammonio-phosphate of magnesium. The residue *C* left upon the filter will consist of uric acid.

β. Aqueous solution. A few drops of the solution *A* are evaporated upon a glass plate, when, should a mere trace only of residue be left, the remainder of this solution may be disregarded, and the calculus considered as free from any appreciable amount of alkaline urates. But should an obvious residue be left, about one-fourth part of the solution may be boiled in a test tube with a little caustic potash, when ammonia, if present, will be given off so as to be recognisable by its odour and by its reactions with test-paper and hydrochloric acid vapour, &c. The remainder of the solution is reduced to a very small bulk by evaporation, treated with strong nitric acid, and evaporated cautiously to dryness, when the production of a pinkish residue, rendered crimson when moistened with ammonia, will be indica-

tive of uric acid. The contents of the capsule are next incinerated, the residue treated with a few drops of water, and the liquid divided into two portions. One is slightly acidified with acetic acid and tested with a drop of oxalate of ammonium, when the production of a white turbidity will indicate the presence of calcium. The other is acidified with hydrochloric acid and evaporated cautiously to dryness, when the production of microscopic cubical crystals will show the presence of sodium.

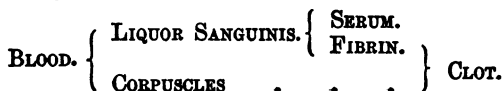
γ. Acid solution. By means of dilute ammonia, the solution *B* is made as nearly neutral as it can be, without having its transparency affected. Acetate of ammonia is then added, the production by which of a white precipitate will indicate the presence of oxalate of calcium, or cystine. The latter body rarely occurs in mixed calculi, and could be readily separated from the oxalate by treatment with ammonia: on evaporating the ammoniacal solution, it would be deposited in the form of hexagonal tablets. To the clear liquid, if no precipitate has formed, or otherwise to the filtrate therefrom, oxalate of ammonium is added in excess, when the deposition of a white precipitate will indicate the presence of calcium, which did not previously exist in the state of an oxalate. Filtration is next performed, if necessary, and an excess of ammonia added to the clear liquid, when the production of a white crystalline precipitate, after stirring for a little while, will prove the presence of phosphoric acid and of magnesium. Should there be no obvious precipitate, sulphate of magnesium is to be added, when the presence of phosphoric acid will be indicated by the formation, after brisk stirring, of a white crystalline precipitate.

δ. Insoluble residue. The residue *C* is to be treated with concentrated nitric acid, and the whole evaporated to dryness, whereby a pink mass will be left, which held over the vapour of ammonia will become crimson, and if subsequently moistened with potash will become purple, reactions characterising uric acid.

§ VII.—BLOOD.

(173.) COAGULATION.

α. Blood as existing in the vessels is seen to consist of red corpuscles floating in a clear liquid, termed the liquor sanguinis. When removed from the living vessels it speedily separates into two portions, a clear yellow liquid, the serum, and a solid red mass, the clot. The liquor sanguinis consists of fibrin and serum; the clot, of fibrin and corpuscles, as seen in the following diagram:—



Thus the chemical investigation of the blood naturally divides itself into separate examinations of the clot and serum.

β. The coagulation of blood is due to the solidification of fibrin, which entangles in its meshes the corpuscles and a considerable portion of the serum, so as to form a firm jelly-like mass. While the blood is circulating through the vessels of living animals, the fibrin exists in a state of perfect solution. The circumstances which determine this state of solution are not well understood; but intimate contact with the living tissues appears to be one very important condition. Out of the body the fibrin speedily solidifies, the coagulation, which is accompanied by a slight evolution of ammonia, being generally complete in about ten minutes' time. Variations of temperature, movement, and exposure to air, modify but never prevent the coagulation. Where the fibrin exists in large quantity, the coagulation takes place more slowly, but the coagulum is firmer and more compact. When blood is removed from persons suffering from an inflammatory condition of system, or when it contains an excess of fibrin, or a deficiency of corpuscles, or when it is collected in a deep narrow vessel, or when its coagulation is retarded by any means, the corpuscles sinking before the coagulation is complete,

exist principally in the lower portion of the clot, while the upper layer consists of nearly colourless fibrin. This colourless layer is termed the *buffy coat*; it is extremely tenacious, and frequently by its slow contraction draws up the edges of the clot, so as to form a cup-like depression.

(174.) FIBRIN.

a. From liquid blood. Fibrin is most easily procured from this source. The blood, before it has had time to coagulate, is rapidly whipped with a few twigs of wood, or well shaken in a bottle with two or three irregular pieces of lead. In this way the fibrin separates more or less completely from the corpuscles, and adheres to the twigs or pieces of lead in the form of loose fibrous masses. These are to be well washed with water, and also with ether, when it is desired to remove the adherent fat.

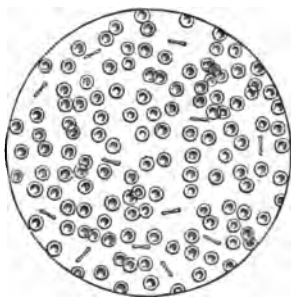
β. From the clot. The preparation of fibrin from coagulated blood is rather more tedious. The clot should be placed upon a cloth, thoroughly broken up by the hand, and washed under a stream of water; when, by alternate washing and kneading, the serum and colouring matter of the clot will pass through the cloth, and a residue of tolerably white fibrin be left thereon.

γ. Properties of fibrin. Fibrin possesses all the chemical properties of coagulated albumen (vide par. 176 *γ*). When examined microscopically it is seen to differ from coagulated albumen in manifesting an organised structure, though of the lowest type, viz., the simply fibrous. This fibrillated arrangement is best seen in the buffy layer of inflammatory blood. When moist fibrin, especially that obtained from the clot, is covered with water rendered *faintly* alkaline by soda, and left at rest for some days in a tolerably warm situation, the greater part of it dissolves, and albumen may be detected in the filtered liquid by the action of heat and nitric acid. Fibrin constitutes about 0.25 per cent. of normal blood.

(175.) CORPUSCLES.

a. Their appearance. When a drop of uncoagulated blood, or a drop of the deep red-coloured serum squeezed out of the clot, is examined under a good quarter-inch object glass with a high eye-piece, the field of the microscope is seen covered with minute coloured cells, of uniform size, circular outline, and non-granular structure, as shown in fig. 68. According to the focussing the edges will appear dark and the centre transparent, or *vice versâ*. Some of the globules may be seen lying upon their edges, some of them adhering to one another by their flat surfaces, forming rouleaus. In the case of the previously uncoagulated blood, a delicate net-work of fibrin will speedily appear. Blood corpuscles appear to consist of a transparent membrane containing a red-coloured fluid. The phenomena of *osmose* may be readily seen under the microscope: thus if a concentrated solution of sulphate of sodium be added, the corpuscles become distorted, their edges uneven, and their dark centres more prominent; if, however, water be added, the corpuscles swell up, their dark centres and defined margins gradually disappear, and finally the cells burst with discharge of their contents.

Fig. 68.



In addition to the above-described red corpuscles, there may generally be seen a few of the colourless or lymph corpuscles. In healthy blood, these exist in a variable but very small proportion compared with the red, than which they are rather larger in size, and less uniform in outline. Moreover, they manifest a faintly granular structure.

β. Their separation. If blood as it is flowing be received into a saturated solution of sulphate of sodium, all

coagulation will be prevented, and by repose the corpuscles will form a bright scarlet layer at the bottom of the vessel. The supernatant fluid may be poured off, and the sediment collected in a filter, and washed with a solution of sulphate of sodium. Or the red liquor, from which the fibrin has been removed by agitation, may be allowed to subside; or the clot may be broken up, well shaken with the serum, and the red fluid so formed be allowed to subside. From either of these fluids a gradual but incomplete deposition of the corpuscles will take place. The supernatant serum may then be poured away, and replaced by a solution of sulphate of sodium, when the corpuscles will behave as in the first instance, and may be collected upon a filter and washed with sulphate of sodium as before.

γ. Red colouring matter. Hæmatosine, or the red colouring matter of blood, is remarkable for the amount of iron which it contains. The ash of blood corpuscles yields fully 30 per cent. of peroxide of iron. The presence of iron in any of the other tissues or fluids, with the exception of the chyle, appears to be due to an admixture of blood. It is possible indeed to obtain a modified hæmatosine free from iron; but no inference can be drawn from the experiment. The chemical reactions of the colouring matter may be recognised by throwing the corpuscles, well washed with sulphate of sodium and drained, into a considerable excess of cold water, when the cell walls burst by endosmosis, and the coloured contents of the cells dissolve in the water, forming a deep-red solution, which by filtration may be made perfectly bright. This red colouring matter is unaffected by ammonia, and is entirely destroyed by ebullition, with the formation of a dirty-coloured coagulum, which dissolves in caustic potash with an indistinct greenish colour.

δ. Hæmatocrystallin, or blood crystals. The formation of these crystals was first discovered by Dr. Otto Funke, of Leipsic. It appears that by the bursting of the red corpuscles as above described an aqueous solution of their contents is obtained, which by very slow evaporation yields crystals

having very definite forms. The blood of some of the lower animals, particularly of the rodents, yields these crystals with great facility, but their production from human blood is always an uncertain operation. A drop or so of the deeply coloured liquid from a portion of clot a day or two old, may be squeezed on to a glass slide and diluted with less than its own bulk of proof spirit. Upon covering the whole loosely with a piece of microscopic glass and setting it aside in a light but not too warm situation, flattened, prismatic red-coloured crystals will sometimes make their appearance in the course of a few hours.

(176.) BLOOD STAINS.

These occasionally form important objects of medico-legal enquiry. The chemical evidence has reference to the colouring matter of the blood; the microscopical to the form of the globules, and the production of hæmine crystals. Recent blood stains are of a bright red colour; older stains of a reddish brown: when on linen or other stuffs, the fibre becomes more or less stiffened.

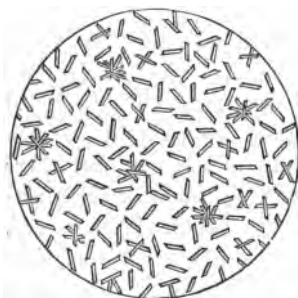
a. Reactions of colour. Supposing the stain to be on some stuff, a strip of the stained portion is cut off, and suspended by means of a thread in a test tube containing a little distilled water. Gradually, streaks of colouring matter will be seen descending from the stuff to the bottom of the tube, and there forming a layer, of a deep-red if the stain be recent, or a reddish-brown colour if the stain be of an older date. If necessary, several of the stained strips may be thus successively treated in the same portion of water, until a sufficiently dark solution is obtained. Stains on knives and other articles are likewise to be treated with cold water, so as to obtain a solution of the colouring matter. To this red or reddish-brown solution a little dilute ammonia is added, whereby the colour will be unaltered or simply brightened, but not changed to a green or crimson. When the solution of the colouring matter is boiled, its colour is entirely destroyed, and a dirty brown precipitate produced. Every stain

forming a reddish solution with cold water, which is unaffected by dilute ammonia, but has its colour destroyed by ebullition, with the formation of a precipitate, is due to blood.

β . Form of the corpuscles. A little of the stained fibre, or, if possible, a little of the dried stain scraped away from the article under examination, is placed upon a glass slide and moistened with a solution of sugar (syrup diluted with about twice its bulk of water) or of pure glycerine reduced to the sp. gr. of 1028. After some time a red coloured liquid will be obtained, which may be covered with a piece of microscopic glass, and examined by a good quarter-inch object-glass. Should the stain be due to blood, the corpuscles, with their characteristic appearances, may in this way be readily detected. In the event of their appearing shrivelled, the addition of a drop or two of water will cause them to expand.

γ . Hæmine crystals. The dry blood stain is extracted with a little glacial acetic acid, and the resulting liquid evaporated at a

Fig. 69.



very gentle heat. In the event of no crystals being thus obtained, a minute quantity of common salt may be added to the dry residue, and the moistening with glacial acetic acid and gentle evaporation repeated. In some cases the moistening with acid has to be performed a third time, before the characteristic dark-coloured rhombic crystals, often arranged in stellate groups, as shown in fig. 69, make their appearance.

The formation of the crystals seems to be facilitated by moistening the residue with water and evaporating therewith, after each evaporation with acetic acid.

(177.) SERUM.

α . General characters. The serum is a pale yellow, transparent, somewhat viscid fluid, having a specific gravity of

about 1030, while that of blood averages about 1055. It has a faintly alkaline reaction, and consists of water holding in solution albumen, fat, certain ill-defined extractive matters, and the inorganic salts. When serum is evaporated to dryness in a water-bath, its aqueous portion is driven off, and a hard, nearly transparent, horny residue left behind. Water constitutes about 80 per cent. of normal blood.

β. Separation of albumen. For this purpose, either of the following methods may be adopted :—*a.* The serum is put into a small capsule, an equal bulk of coarsely powdered crystals of sulphate of sodium added to it, and the whole boiled until complete coagulation takes place, when, on filtering the boiling liquid, a perfectly clear and nearly colourless solution, quite free from albumen, will rapidly pass through. This method is equally applicable to serum containing any amount of colouring matter, and even to the broken-up clot itself. *b.* The serum is made neutral or *very faintly* acid with acetic acid, boiled and filtered. By this means the whole of the albumen will coagulate in flakes and remain on the filtering paper, while a clear liquid, termed the *serosity*, will pass through. The precipitate of albumen is to be washed with hot water, and dried in a water-bath; moreover, a minute amount of earthy salts may be removed from it by boiling dilute hydrochloric acid, as also a small proportion of fat by boiling alcohol.

γ. Properties of albumen. Albumen, as it exists in the blood and other animal fluids, is in a state of solution, but is capable of being coagulated by heat. The temperature at which coagulation takes place varies with the alkalinity of the fluid, and with the amount of albumen present. Serum of blood coagulates at the temperature of about 160° Fahr. Albumen once coagulated cannot again be obtained in the form of a solution coagulable by heat. Normal blood contains on the average about 7 per cent. of dissolved albumen. Fibrin and coagulated albumen agree in the following characters: they are insoluble in water, alcohol, and ether; but are soluble in potash, from which solution they are

reprecipitated by neutralisation with an acid. If to the potash solution acetic acid be freely added, the albumen will be at first precipitated, but subsequently redissolved in the excess of acid. Moreover, coagulated albumen and fibrin are soluble, though with difficulty, in acetic acid. Albumen and fibrin dissolve in boiling hydrochloric acid, forming deep purple solutions.

When albumen or fibrin is heated upon platinum foil, a minute white ash, consisting principally of phosphate of calcium, remains. This proportion of earthy phosphate appears to be an integral constituent of the albuminous principles. If dried albumen or fibrin be heated in a reduction tube, into the mouth of which there have been inserted a piece of red litmus and a piece of lead paper, the red litmus will become blue and the lead paper black, reactions indicating respectively the presence of nitrogen and sulphur.

d. *Fat of blood.* The condition in which fat exists in the serum is not well understood. Some portion of it is precipitated with the albumen, the remainder being dissolved in the serosity. As a rule, serum is perfectly bright, and fat globules cannot be detected in it by microscopical examination; yet the fat being soluble in ether, does not appear to be saponified. In order to extract the fat, the dried residue left by the evaporation of serum upon a water-bath is pulverised, and the powder agitated for some time, with three or four times its bulk of ether; the whole is allowed to subside, and after some hours the ether poured off and evaporated to dryness on a water-bath, when a small quantity of a yellow semi-solid fat will be left. This may be treated with cold alcohol, when a crystalline fat will be dissolved and an oily fat be left unacted upon.

e. *Extractives and salts.* The serosity consists of an aqueous solution of certain ill-defined organic compounds, known as extractives, and of the usual alkaline salts. All the organic constituents of the serum, with the exception of the albumen and fat, receive the name of extractives. If the serosity be carefully evaporated almost to dryness, very beautiful cubes of common salt

crystallise out. Moreover, sulphuric and phosphoric acids can readily be detected; the former by the production of a white precipitate with nitrate of barium and nitric acid; the latter by the addition of sulphate of magnesium, ammonia, and chloride of ammonium, when on stirring a white crystalline precipitate is thrown down. If the serosity be evaporated to dryness, and ignited, a white fusible ash will remain, which may be examined according to the directions for the analysis of animal ashes, and will be found to contain chlorine, carbonic, phosphoric, and sulphuric acids, with sodium, potassium, and traces of calcium.

ζ. Serum containing urea. It is probable that the blood always contains minute traces of urea; while in certain forms of disease, particularly in Bright's disease, its amount becomes very sensible. In order to detect it, the albumen is removed from the serum by either of the methods described in par. β, and the serosity evaporated carefully to dryness. If the process *b* be adopted, and the chloride of sodium in the dried residue be seen to crystallise in well-marked octahedrons instead of cubes, the presence of urea is tolerably certain. In any case the dry residue is warmed with a little strong alcohol and filtered, the alcoholic filtrate evaporated to dryness, the residue dissolved in a very small quantity of distilled water, and the aqueous solution filtered and concentrated in a watch-glass. To the cold concentrated liquid, an equal bulk of cold colourless nitric acid is next added, when, if urea be present, a crystalline deposit of nitrate of urea will be produced, which can be examined microscopically. The production of a crystalline deposit on the addition of nitric acid is in itself almost conclusive as to the presence of urea.

η. Serum containing bile. In jaundice, the serum of the blood is of a much darker yellow colour than usual, owing to the presence of biliary colouring matter, which may be identified by adding to the serum a little nitric acid, when the albumen will be precipitated of a bluish or greenish colour: or some of the serum, just neutralised with acetic acid, may be precipitated with excess of alcohol, the alcoholic solution evaporated to a small bulk

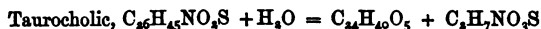
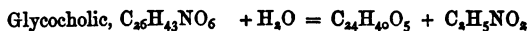
in a water-bath, and residue examined by Pettenkofer's test, and by the nitric acid test (par. 153 β).

θ . Serum containing sugar. Sugar exists abundantly in the blood, in cases of diabetes. Recent researches have moreover shown that the *post-mortem* blood of the inferior cava, and of the right side of the heart, habitually contains sugar in very definite amount. In order to detect it, the albumen of the blood must be removed by means of sulphate of sodium, as in par. 177 β , and to the clear filtrate a drop or two of a solution of sulphate of copper, and then an excess of caustic potash be added, whereby a deep blue-coloured liquid will be produced, which, on the application of heat, will deposit a red or orange-yellow coloured precipitate of suboxide of copper. The alkaline solution of tartrate of copper described in par. 162 δ . may be conveniently substituted for the mixture of sulphate of copper and caustic potash.

§ VIII.—MISCELLANEOUS ANIMAL PRODUCTS.

(178.) BILE.

α . Composition. Human bile contains the sodium salt of a peculiar acid known as glycocholic acid, having a similar constitution to the glycobenzoic or hippuric acid found in urine. It also contains in small quantity the sodium salt of taurocholic acid. These glycocholates and taurocholates, when boiled with dilute hydrochloric acid, absorb water, and break up into cholic acid and glycocine and taurine respectively, thus:—



The bile also contains water, mucus, cholesterine, fat and colouring matter, the reactions of which last have been described under the head of biliary urine (par. 153).

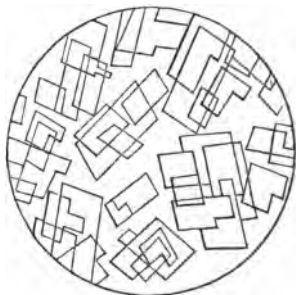
β . Detection of cholic acid. This is effected by Pettenkofer's test, which is performed in the following manner: To a little diluted bile, or any liquid containing bile, rather more

than half its bulk of strong sulphuric acid is added very gradually, the tube in which the mixture is made being kept cool by immersion in water. A minute quantity of powdered white sugar, or its equivalent of syrup, is then introduced, the liquid well agitated and mixed with more sulphuric acid. By this means the temperature is gradually raised to the requisite point, when a deep purplish-crimson colour makes its appearance. Very minute quantities of bile may be detected by evaporating any suspected liquid with a drop of sulphuric acid and a decigrain of sugar in a water-bath.

γ. Biliary calculi. Concretions of variable appearance are occasionally formed in the gall-bladder, sometimes in very considerable numbers. When numerous, they are of about the size of peas, and have an irregular angular shape, with flattened sides. When solitary, they are usually of larger bulk and of an oval form. They have a soapy feel, a fawn-yellow colour, and are easily crushed by pressure. They consist principally of *cholesterine* and an insoluble combination of *bile pigment with lime*.

When rich in cholesterine, they float upon water. To detect cholesterine, the powdered calculus is boiled in alcohol, and the solution filtered, when, on cooling, colourless transparent plates crystallise out of the yellow liquid. These crystals, when examined microscopically, are seen to consist of delicate, nearly square, rhombic plates, superimposed one upon the other as shown in fig. 70. Cholesterine

Fig. 70.

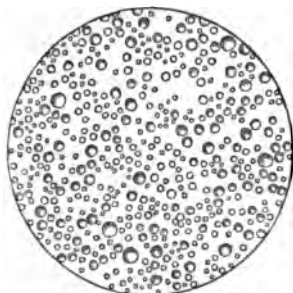


$C_{26}H_{44}O$, is soluble in ether and hot alcohol, very slightly soluble in cold alcohol, not at all soluble in water. It seems to be a species of solid alcohol.

(179.) MILK.

This secretion consists of water holding in solution casein, lactine, and salts, and holding in suspension an abundance of

Fig. 71.



fat globules, to the presence of which the opaque white appearance of milk is due. Normal milk has an alkaline reaction to test paper. The specific gravity of human milk averages about 1030.

a. Fat globules. These may be easily recognised under the microscope: they are of various sizes, and have well-defined dark margins, as shown in fig. 71. They appear to be surrounded by delicate cell-

membranes, as they cannot be made to unite by pressure until after the addition of a little acetic acid. The milk secreted soon after delivery contains large, granular, fatty corpuscles, known as colostrum corpuscles. In milk abscess &c., pus and blood globules may be occasionally detected.

β. Casein. When ordinary or skimmed milk is evaporated, a scum forms on its surface, which, if removed, is soon replaced by a fresh one, and so on repeatedly. This property of forming a scum on evaporation was formerly considered peculiar to fluids containing casein. Casein differs from the other albuminous bodies in being coagulable not by heat, but by the addition of a little very dilute acid, or by contact with decomposing animal membrane. Cheese is casein which has been precipitated by rennet, the dried decomposing lining membrane of the stomach of the calf. When skimmed milk is rendered slightly acid with acetic acid, and gently warmed, the casein coagulates, and may be collected on a filter, washed with hot water, and subsequently with hot alcohol: it manifests all the usual properties of the albuminous bodies.

γ. *Lactine*. Sugar of milk, or lactine, may be readily detected in whey which has been separated by filtration from the coagulated casein. On adding a drop or two of sulphate of copper, and then an excess of potash, a red precipitate of sub-oxide of copper will be produced on boiling the mixture. Sugar of milk may be obtained crystalline by careful evaporation of the whey. It does not readily undergo the alcoholic fermentation; but by the action of putrefying curd becomes rapidly converted into lactic acid.

δ. *Mineral salts*. The ash of milk contains the same constituents as most animal ashes; the relative proportion of earthy phosphates is very large, and potassium exists in it to a greater extent than sodium.

(18c.) BONE.

Bone consists principally of phosphate of calcium, deposited in an animal basis. When bone is soaked for some time in dilute hydrochloric acid, its earthy matter is dissolved out, and a flexible elastic mass, having the exact form of the original bone, is left unacted upon. This residue consists of gelatine, which by long boiling dissolves in water, forming a solution that gelatinises on cooling. The solution is precipitated by tannic acid, but not by acetic acid or by ferrocyanide of potassium. Pure gelatine does not contain sulphur. By the incineration of bone, its animal matter is burnt off, and a brittle, white, earthy residue, having the exact form of the original bone, remains. This earthy residue consists principally of phosphate of calcium, with a little carbonate of calcium and phosphate of magnesium, also minute quantities of fluoride of calcium; which latter substance may be detected more readily, however, in fossil bones.

INDEX.

ACE

- A** CETATES, reactions of, 135
 Acid, arsenious, 157
 — cholic, 224
 — hippuric, 192
 — hydrochloric, toxicology of, 144
 — hydrofluosilicic, as a reagent, 85
 — nitric, toxicology of, 142
 — oxalic, „ „ 147
 — prussic, „ „ 173
 — sulphuric, as a reagent, { 88
 { 93
 — — toxicology of, 139
 — tartaric, as a reagent, 68
 — uric, 190
 Acidity, criterion of, 6
 Acids and bases, 52
 — anhydrous, 11
 — binary, 6
 — dibasic, 7, 8
 — monobasic, 7
 — reactions of, 107
 — solutions to be tested for, 95
 — table for detection of, 96
 — — preliminary testing, 90
 — ternary, 6
 — tribasic, 8
 Adapters, 37
 Albumen from blood, 220
 Albuminous urine, 195
 Alkaline solutions, reactions of, 110
 Alkaloidal salts, 9
 Aluminum, reactions of, 122
 Ammonia, as a precipitant, 74
 — — — reagent, 79
 — — — solvent, 80
 Ammonium, carbonate of, as a pre-
 cipitant, 83
 — oxalate of, as a reagent, 85

ATO

- Ammonium, phosphate of, as a re-
 agent, 68
 — salts of, 9
 — sulphide of, as a precipitant, 73
 — — as a solvent, 70
 Analysis, course of operations in,
 52
 Anhydrides, 11
 Anhydrous acids, 11
 Animal matter, ashes of, 184-186
 — — components of, 183
 — — products, various, 224
 Antimonial salts, reactions of, 169
 Antimony in organic mixtures, 173
 — Marsh's test for, 171
 — reactions of, 113, 169
 — Reinsch's test for, 172
 — toxicology of, 169
 — — in solution, 170
 Apparatus, cleaning glass, 39
 — connections of, 22
 — supports for, 32
 Arsenic, Marsh's test for, 161
 — — original process, 162
 — — modified process, 164
 — metallic ring, 158
 — reactions of, 112
 — Reinsch's test for, 165
 — — — impediments to, 167
 — — various forms of, 168
 Arsenious acid, reactions of, 157
 — — toxicology of, 157
 — — — crystals of, 157
 — — — dissolved, 159
 — — — reduction of, 159
 — — — in organic mixtures, 169
 Ashes of animal matter, 184, 186
 Atomic weights, 2

BAR

- BARIUM**, chloride of, as a reagent, 98
 — reactions of, 123
 Bases and acids, 52
 — classification of, 64
 — first group of, recognised, 65
 — — — table of, 68
 — second group of, recognised, 73
 — — — table of, 76
 — third group of, recognised, 82
 — — — table of, 83
 Baths, sand, 33
 — water, 34
 Beads, borax, 60
 Benzoates, reactions of, 135
 Biliary calculi, 224
 — urine, 199
 Bile, 224
 — from blood, 223
 Binary acids, 6
 Bink's burette, 45
 Bismuth, reactions of, 113
 — solutions, 72
 Blood, albumen from, 220
 — bile in, 223
 — coagulation of, 215
 — colouring matter of, 218
 — composition of, 215
 — corpuscles, 217
 — crystals, 218
 — extractives and salts of, 222
 — fat of, 222
 — fibrin from, 216
 — in urine, 205
 — serum from, 220
 — stains, reactions of, 219
 — sugar in, 224
 — urea in, 222
 Blowing glass bulbs, 21
 Blowpipe, Black's, 16
 — brazier's, 16
 — examination, 53
 — flame, colouration of, 56
 — flames, 17
 — Herapath's gas, 15
 — incrustations, 59
 — the use of, 17
 Bone, 226
 Borates, reaction of, 135
 Borax beads, 60
 Boring corks, 23

CON

- Bottle, drop, 24
 — wash, 25
 Bromides, reactions of, 131
 Bulbs, blowing, 21
 Burette, Bink's, 45
 Burners, charcoal, 14
 — gas, 14
CADMIUM, reactions of, 117
 Calcium, chloride of, as a reagent, 198
 — oxalate of, urinary deposit of, 202
 — — calculus, 209
 — reactions of, 124
 Calculi, action of heat on, 209
 — analysis (systematic) of, 213
 — biliary, 225
 — construction of, 207
 — cystine, 209
 — earthy phosphate, 208
 — — — reactions of, 212
 — examination of, 210
 — ignition of, 210
 — mulberry, 208
 — oxalate of calcium, 209
 — — — reactions of, 212
 — pulverising, 210
 — solution of, 213
 — tests, special for, 211
 — uric acid, 208
 — — — reactions of, 212
 Carbonates, reactions of, 119
 Casein, 226
 Charcoal burners, 13
 Chemical equations, 6
 — formulæ, 5
 — urinary deposits, 201
 Chlorates, reactions of, 127
 Chlorides, formulæ of, 4
 — reactions of, 130
 Cholic acid, 224
 Chromates, reactions of, 126
 Chromium, reactions of, 122
 Chylous urine, 200
 Cleaning-glass apparatus, 39
 Cobalt, nitrate of, ignitions with, 57
 — reactions of, 118
 Colouring-matter of blood, 218
 Combining proportions, 1
 Complex salts, 9
 Condenser, Liebig's, 38

CON

- Connections of apparatus, 22
 Copper, reactions of, 116, 155
 — solutions, 73
 — toxicology of, 155
 — — dissolved, 155
 — — in organic mixtures, 156
 Corks, boring, 23
 Corpuscles, blood, 217
 Corrosive sublimate, reactions of,
 149
 — toxicology of, 149
 — — solid, 149
 — — dissolved, 150
 — — crystals of, 150
 — — in organic mixtures, 151
 Counterpoises, 46
 Crucibles, heating, 34
 Crystals, blood, 218
 — hæmine, 220
 Cyanides, reactions of, 132
 Cystine, 203
 — calculi, 209

DASH'D SYMBOLS, 5

- Decantation, 32
 Decimal weights, 46
 Deposits, urinary, 200
 — — chemical, 201
 — — organised, 203
 Desiccation, 39, 46
 Dibasic acids, 6
 Distillation, 37
 Drop-bottle, 24
 Drying apparatus, 39
 — precipitates, 40

EARTHY SALTS, reactions of, 105

- Ebullition, 35
 Elementary bodies, 1
 Elements, table of, 3
 Equations, chemical, 6
 Equivalents, 1
 Evaporation, 35

FAT of blood, 222

- Fatty urine, 199
 Ferric salts, reactions of, 120
 Ferrous salts, reactions of, 120

HYD

- Fibrin, 216
 Filters, double, 30
 Filters, making, 29
 — plain, 30
 — ribbed, 29
 Filtration, 28
 Flame, blowpipe, 17
 — — colouration of, 56
 — — oxidising, 17
 — — reducing, 17
 Flasks, measure, 43
 — specific gravity, 48
 Fluorides, reactions of, 136
 Formulæ, chemical, 5
 — of chlorides, 4
 — of hydrides, 3
 — of salts, rational, 10
 Funnel tubes, 26
 Fusible salts, 56
 Fusions, 41

GALLON, decimal division of, 43

- Gas blowpipe, Herapath's, 15
 — burners, 14
 — evolving tubes, 27
 — washing tubes, 27
 Glass bending, 19
 — tube and rod, cutting, 18
 — vessels, heating, 35
 — working, 18
 Graduated measures, 44
 — pipettes, 44
 Gravity, specific, 47

HÆMATO-CRYSTALLINE, 218

- Hæmine crystals, 220
 Heat, action of, on calculi, 209
 Heating glass vessels, 35
 — crucibles, 34
 Hippuric acid, 192
 Hydracids, 7
 Hydrides, formulæ of, 3
 Hydrochloric acid, as a precipitant,
 65
 — — reactions of, 144
 — — toxicology of, 144
 — — concentrated, 144
 — — diluted, 145
 — — in organic mixtures, 146

HYD

Hydrofluosilicic acid as a reagent, 85
 Hydrometer, 49

IGNITIONS, 41

— with nitrate of cobalt, 57
 Incrustations, blowpipe, 59
 Infusible salts, 57
 Insoluble compounds, in acids, 102
 — oxalates, 149
 Iodides, reactions of, 131
 Iron, perchloride of, as a reagent, 100
 — reactions of, 120

JACKET STANDS, 33

KIESTEIN URINE, 200

LACTINE, 226

Lead solutions, 72
 — reactions of, 115, 133
 — toxicology of, 153
 — — — solid, 153
 — — — dissolved, 154
 — — — in organic mixtures, 154
 Liebig's condenser, 38
 Liquids, analysis of, 109
 — specific gravity of, 48
 Lithates, urinary deposits of, 201
 Lutes, 24

MAGNESIUM, reactions of, 124

Manganese, reactions of, 119
 Marsh's antimony test, 171
 — arsenic test, 161
 — — — original process, 162
 — — — modified process, 164
 Measure flasks, 44
 Measures, graduated, 45
 Measuring, 44
 Mercuric salts, reactions of, 115
 Mercurous salts, reactions of, 115
 Mercury, dichloride of, toxicology of, 149
 — reactions of, 114
 — solutions, 72

PLA

Milk, 226
 — casein of, 226
 — fat globules of, 226
 — lactine of, 226
 — mineral salts of, 226
 Molecular weights, 2
 Monobasic acids, 7
 Morphia in solution, 180
 — toxicology of, 179
 Mucus from urine, 204
 Mulberry calculus, 208
 Multivalent salts, 8

NEUTRALISATION, 74

Nickel, reactions of, 118
 Nitrates, reactions of, 127, 143
 Nitric acid, toxicology of, 142
 — — — concentrated, 142
 — — — diluted, 143
 — — — in organic mixtures, 144
 Notation, symbolic, 5

OPIATE LIQUIDS, 180

Organic constituents, ultimate, 184
 Oxacids, 6
 Oxalate of calcium, urinary deposit of, 202
 Oxalates, reactions of, 133
 — insoluble, 105, 149
 Oxalic acid, reactions of, 147, 212
 — toxicology of, 147
 — — — dissolved, 147
 — — — in organic mixtures, 148
 — — — insoluble form, 149
 — — — solid, 147
 Oxides, reactions of, 106
 Oxidising blowpipe flame, 16

PASTILLES, 18

Phosphates, reactions of, 132
 — earthy, urinary deposits of, 201
 Pipettes, delivery, 44
 — graduated, 45
 — making, 22
 Platinum, perchloride of, as a reagent, 86

POT

- Potash, as a reagent, 71, 78
 — solutions, 81
 Potassium, sulphate of, as a reagent, 84
 Powders, specific gravity of, 51
 Precipitant, carbonate of ammonium as a, 83
 — hydrochloric acid as a, 65
 — sulphide of ammonium as a, 73
 — sulphuretted hydrogen as a, 69
 Precipitates, aspects of, 63
 — drying, 40
 — formation of, 62
 — removal of, 32
 — washing, 31
 Precipitation by ammonia, 74
 — by water, 66
 Proportions, combining, 1
 Prussic acid, toxicology of, 173
 Purpurine, 193
 Pus from urine, 204

REACTIONS of acetates, 135

- — acids, 107
 — — alkaline solutions, 110
 — — aluminum, 122
 — — ammonium, 126
 — — antimony, 113, 169
 — — arsenic, 112, 157
 — — barium, 121
 — — benzoates, 135
 — — bismuth, 113
 — — blood stains, 219
 — — borates, 135
 — — bromides, 131
 — — cadmium, 117
 — — calcium, 124
 — — carbonates, 128
 — — chlorates, 127
 — — chlorides, 130
 — — chromates, 126
 — — chromium, 122
 — — cobalt, 118
 — — compounds in solution, 109
 — — copper, 116, 155
 — — corrosive sublimate, 149
 — — cyanides, 132
 — — earthy salts, 105
 — — ferric salts, 120
 — — ferrous salts, 120

REA

- Reactions of fluorides, 106
 — — hippuric acid, 193
 — — hydrochloric acid, 144
 — — iodides, 131
 — — iron, 120
 — — lead, 115, 153
 — — liquid substances, 109
 — — magnesium, 124
 — — manganese, 119
 — — mercuric salts, 115
 — — mercurous salts, 115
 — — mercury, 114
 — — miscellaneous salts, 108
 — — morphia, 179
 — — nickel, 118
 — — nitrates, 127, 143
 — — nitric acid, 143
 — — oxalates, 133
 — — oxalic acid, 147
 — — — — calculi, 212
 — — oxides, 106
 — — phosphates, 132
 — — phosphatic calculi, 212
 — — potassium, 125
 — — prussic acid, 173
 — — silicates, 137
 — — silver, 116
 — — sodium, 126
 — — stannic salts, 112
 — — stannous salts, 111
 — — strontium, 124
 — — strychnia, 175
 — — sulphates, 130
 — — sulphides, 106, 129
 — — sulphites, 130
 — — sulphuric acid, 139
 — — sulphhydrates, 129
 — — tartrates, 134
 — — tin, 111
 — — urea, 189
 — — uric acid, 190
 — — — — calculi, 212
 — — zinc, 121
 Reagent, ammonia as a, 79
 — — chloride of barium as a, 98
 — — — — calcium as a, 98
 — — hydrofluosilicic acid as a, 85
 — — nitrate of silver as a, 96
 — — oxalate of ammonium as a, 85
 — — perchloride of iron as a, 100

REA

- Reagent, perchloride of platinum as a, 86
 — phosphate of ammonium as a, 68
 — potash as a, 71, 78
 — sulphate of potassium as a, 84
 — sulphuric acid as a, 88, 93
 — tartaric acid as a, 86
 Reagents, addition of, 63
 Reducing blowpipe flame, 17
 Reduction tubes, 18
 Reinsch's test for antimony, 172
 — — — arsenic, 165
 — — — impediments to, 167
 Retorts, tube, 25

SACCHARINE URINE, tests for, 196

- Salts, alkaloidal, 9
 — ammonium, 9
 — complex, 9
 — double decomposition of, 12
 — earthy, nature of, 75
 — — reactions of, 105
 — fusible, 56
 — infusible, 57
 — mineral, of milk, 226
 — — of urine, 194
 — miscellaneous, 108
 — multiequivalent, 8
 — rational formulæ of, 11
 — reducible, 58
 — simple, 8
 — volatile, 55

Sand-baths, 34

Sealed tube making, 21

Sealing test tubes, 20

Sediment from normal urine, 203

Serum of blood, 216

Silicates, reactions of, 137

Silver, nitrate of, as a reagent, 96

— reactions of, 116

— solutions, 73

Simple salts, 9

Solids, specific gravity of, 50

Solutions made, 61

Solvent, ammonia as a, 80

— sulphide of ammonium as a, 70

Solvents, addition of, 63

Specific gravity, 47

— — flasks, 48

TOX

Specific gravity of liquids, 48

— — of powders, 51

— — of solids, 50

Stains, blood, reactions of, 219

Stands, jacket, 33

— tripod, 33

Stannic salts, reactions of, 112

Stannous salts, reactions of, 111

Stirring rods, 18

Strontium, reactions of, 124

Strychnia, appearance of, 176

— colour tests, 177

— in organic mixtures, 178

— toxicology of, 175

Subliming tubes, 18

Sugar in blood, 224

Sulphates, reactions of, 130

Sulphides, reactions of, 106, 129

Sulphites, reactions of, 130

Sulphuretted hydrogen as a precipitant, 65

— — bulb, 27

Sulphuric acid, as a reagent, 93

— — reactions of, 139

— — toxicology of, 139

— — concentrated, 139

— — diluted, 140

— — — in organic mixtures, 141

— — — stains on clothing, 141

Sulphydrates, reactions of, 129

Supports for apparatus, 33

Symbolic notation, 5

Symbols, dash'd, 5

Synoptic formulæ, 11

TABLE for detection of acids, 96

— — — of bases of group one, 68

— — — — of group two, 76

— — — — of group three, 84

— of blowpipe reactions, 55

— of elements, 3

— of preliminary testing for acids, 90

Tartrates, reactions of, 134

Ternary acids, 6

Test tubes, sealing, 20

Tin, reactions of, 111

Tissues, composition of, 183

Toxicological examinations, 139

TOX

- Toxicology of antimony, 169
 — — — arsenious acid, 157
 — — — copper, 155
 — — — corrosive sublimate, 149
 — — — hydrochloric acid, 144
 — — — hydrocyanic acid, 173
 — — — lead, 153
 — — — mercury, 149
 — — — morphia, 179
 — — — nitrates, 143
 — — — nitric acid, 142
 — — — oxalic acid, 147
 — — — prussic acid, 173
 — — — strychnia, 175
 — — — sulphuric acid, 139
 Tripod stands, 34
 Tube retorts, 25
 Tubes, funnel, 26
 — gas-evolving, 26
 — joined, 23
 — subliming, 18
 — reduction, 18
 Tubing, vulcanite, 22

URATES, urinary deposits of, 201

- Urea, crystals of, 190
 — from blood, 222
 — its detection, 189
 — preparation of, 189
 — properties of, 190
 Uric acid, crystalline form of, 192
 — — its detection, 190
 — — preparation of, 191
 — — properties of, 191
 — — urinary deposit of, 201
 — — calculus, 208
 Urinary calculi, 207
 — — construction of, 207
 — deposits, 200
 — — chemical, 201
 — — — of cystine, 203
 — — — earthy phosphates, 201
 — — — lithates, 201
 — — — oxalate of calcium,
 202
 — — — — urates, 201
 — — — — uric acid, 201

ZIN

- Urinary deposits, organised, 203
 — — — of blood, 205
 — — — — mucus, 204
 — — — — normal sediment, 203
 — — — — pus, 204
 Urine, 188
 — abnormal, 195
 — albuminous, 195
 — — tests for, 195
 — appearance of, 205
 — ash of, 194
 — biliary, 199
 — chemical examination of, 206
 — chylous, 200
 — clinical examination of, 205
 — colouring-matters of, 193
 — crystals from evaporation of, 194
 — extractive matters of, 193
 — fatty, 199
 — kiestein, 200
 — mineral salts of, 194
 — normal, properties of, 188
 — quantity of, 206
 — saccharine, 196
 — — tests for, 197
 — specific gravity of, 205

VOLATILE SALTS, 55

- Volatilisations, 42
 Vulcanite tubing, 22

WASH BOTTLE, 25

- Washing precipitates, 31
 — tube, gas, 28
 Water baths, 34
 — precipitations by, 66
 Weighing, 46
 Weights, atomic, 2
 — decimal, 47
 — molecular, 2

ZINC, blowpipe examination of,

- 57
 — reactions of, 121

LONDON: PRINTED BY
SPOTTISWOODE AND CO., NEW-STREET SQUARE
AND PARLIAMENT STREET

[APRIL 1869]

GENERAL LIST OF WORKS

PUBLISHED BY

MESSRS. LONGMANS, GREEN, AND CO.

PATERNOSTER ROW, LONDON.

Historical Works.

LORD MACAULAY'S WORKS. Complete and Uniform Library Edition. Edited by his Sister, Lady TREVELYAN. 8 vols. 8vo. with Portrait, price £5 5s. cloth, or £3 8s. bound in tree-calf by Rivière.

The HISTORY of ENGLAND from the Fall of Wolsey to the Death of Elizabeth. By JAMES ANTHONY FROUDE. M.A. late Fellow of Exeter College, Oxford. VOLS. I. to X. in 8vo. price £7 2s. cloth.

VOLS. I. to IV. the Reign of Henry VIII. Fourth Edition, 54s.

VOLS. V. and VI. the Reigns of Edward VI. and Mary. Third Edition, 28s.

VOLS. VII. and VIII. the Reign of Elizabeth, VOLS. I. and II. Fourth Edition, 28s.

VOLS. IX. and X. the Reign of Elizabeth, VOLS. III. and IV. 32s.

The HISTORY of ENGLAND from the Accession of James II.

Lord MACAULAY.

LIBRARY EDITION, 5 vols. 8vo. £4.

CABINET EDITION, 8 vols. post 8vo. 48s.

PEOPLE'S EDITION, 4 vols. crown 8vo. 16s.

An ESSAY on the HISTORY of the ENGLISH GOVERNMENT and Constitution, from the Reign of Henry VII. to the Present Time. By JOHN EARL RUSSELL. Fourth Edition, revised. Crown 8vo. 6s.

On PARLIAMENTARY GOVERNMENT in ENGLAND: Its Origin, Development, and Practical Operation. By ALPHRUS TODD, Librarian of the Legislative Assembly of Canada. In Two Volumes. VOL. I. 8vo. 16s.

HISTORY of the REFORM BILLS of 1866 and 1867. By HOMERSHAM COX, M.A. Barrister-at-Law. 8vo. 7s. 6d.

Antient Parliamentary Elections, a History shewing how Parliaments were Constituted, and Representatives of the People Elected, in Ancient Times. By the same Author. 8vo. 8s. 6d.

Whig and Tory Administrations during the Last Thirteen Years. By the same Author. 8vo. 5s.

- The HISTORY of ENGLAND** during the Reign of George the Third.
By the Right Hon. W. N. MASSEY. Cabinet Edition. 4 vols. post 8vo. 24s.
- The CONSTITUTIONAL HISTORY of ENGLAND**, since the Accession of George III. 1760—1860. By Sir THOMAS ERSKINE MAY; O.B. Second Edition. 2 vols. 8vo. 33s.
- HISTORICAL STUDIES.** By HERMAN MERIVALE, M.A. 8vo. price 12s. 6d.
- The OXFORD REFORMERS of 1498**; being a History of the Fellowship of John Colet, Erasmus, and Thomas More. By FREDERIC SEEBOHM. 8vo. 12s.
- LECTURES on the HISTORY of ENGLAND**, from the earliest Times to the Death of King Edward II. By WILLIAM LONGMAN. With Maps and Illustrations. 8vo. 15s.
- The HISTORY of the LIFE and TIMES of EDWARD the THIRD.**
By WILLIAM LONGMAN. With 9 Maps, 8 Plates, and 16 Woodcuts. 2 vols. 8vo. 28s.
- HISTORY of CIVILISATION** in England and France, Spain and Scotland. By HENRY THOMAS BUCKLE. Fifth Edition of the entire Work, with a complete INDEX. 3 vols. crown 8vo. 24s.
- WATERLOO LECTURES**: a Study of the Campaign of 1815. By Colonel CHARLES C. CHESNEY, R.E. late Professor of Military Art and History in the Staff College. 8vo. with Map, 10s. 6d.
- HISTORY of GRANT'S CAMPAIGN for the CAPTURE of RICHMOND, 1864—1865**: with an Outline of the Previous Course of the American Civil War. By JOHN CANNON. Post 8vo. 12s. 6d.
- DEMOCRACY in AMERICA.** By ALEXIS DE TOCQUEVILLE. Translated by HENRY REEVE. 2 vols. 8vo. 21s.
- HISTORY of the REFORMATION in EUROPE** in the Time of Calvin. By J. H. MERLE D'AUBIGNÉ, D.D. Vols. I. and II. 8vo. 28s. VOL. III. 12s. VOL. IV. 16s. VOL. V. price 16s.
- HISTORY of FRANCE**, from Clovis and Charlemagne to the Accession of Napoléon III. By EYRE EVANS CROWE. 5 vols. 8vo. £4 13s.
- The HISTORY of GREECE.** By C. THIRLWALL, D.D. Lord Bishop of St. David's. 8 vols. fcp. 8vo. price 28s.
- The TALE of the GREAT PERSIAN WAR**, from the Histories of Herodotus. By GEORGE W. COX, M.A. New Edition. Fcp. 3s. 6d.
- GREEK HISTORY** from Themistocles to Alexander, in a Series of Lives from Plutarch. Revised and arranged by A. H. CLOUGH. Fcp. with 44 Woodcuts, 6s.
- CRITICAL HISTORICAL HISTORY of the LANGUAGE and LITERATURE of Ancient Greece.** By WILLIAM MURE, of Caldwell. 5 vols. 8vo. £3 9s.
- HISTORY of the LITERATURE of ANCIENT GREECE.** By Professor K. O. MÜLLER. Translated by the Right Hon. Sir GEORGE CORNEWALL LEWIS, Bart. and by J. W. DONALDSON, D.D. 3 vols. 8vo. 21s.
- HISTORY of the CITY of ROME** from its Foundation to the Sixteenth Century of the Christian Era. By THOMAS H. DYER, LL.D. 8vo. with 2 Maps, 15s.

HISTORY of the ROMANS under the EMPIRE. By the Rev. C. MERRIVALE, LL.D. 8 vols. post 8vo. 48s.

The FALL of the ROMAN REPUBLIC: a Short History of the Last Century of the Commonwealth. By the same Author. 12mo. 7s. 6d.

The HISTORY of INDIA, from the Earliest Period to the close of Lord Dalhousie's Administration. By JOHN CLARK MARSHMAN. 3 vols. crown 8vo. 22s. 6d.

INDIAN POLITY: a View of the System of Administration in India. By Major GEORGE CHESNEY, Fellow of the University of Calcutta. 8vo. with Map, 21s.

HISTORY of the FRENCH in INDIA, from the Founding of Pondichery in 1674 to its Capture in 1761. By Lieutenant-Colonel G. B. MALLESON, Bengal Staff Corps. 8vo. 16s.

REALITIES of IRISH LIFE. By W. STEUART TRENCH, Land Agent in Ireland to the Marquess of Lansdowne, the Marquess of Bath, and Lord Digby. With Illustrations from Drawings by the Author's Son, J. TOWNSEND TRENCH. Third Edition, with 30 Plates. 8vo. 21s.

JOURNALS, CONVERSATIONS, and ESSAYS relating to IRELAND. By NASSAU WILLIAM SENIOR. 2 vols. post 8vo. 21s.

MODERN IRELAND; its Vital Questions, Secret Societies, and Government. By an ULSTERMAN. Post 8vo. 6s.

IRELAND in 1868 the BATTLE-FIELD for ENGLISH PARTY STRIFE: its Grievances, Real and Factitious; Remedies, Abortive or Mischievous. By GERALD FITZGIBBON. Second Edition. 8vo. 8s. 6d.

AN ILLUSTRATED HISTORY of IRELAND, from the Earliest Period to the Year of Catholic Emancipation. By MARY F. CUSACK. Second Edition, revised and enlarged. 8vo. 18s. 6d.

CRITICAL and HISTORICAL ESSAYS contributed to the *Edinburgh Review*. By the Right Hon. LORD MACAULAY.

LIBRARY EDITION, 3 vols. 8vo. 36s.

CABINET EDITION, 4 vols. post 8vo. 24s.

TRAVELLER'S EDITION, in One Volume, square crown 8vo. 21s.

PEOPLE'S EDITION, 2 vols. crown 8vo. 8s.

GOD in HISTORY; or, the Progress of Man's Faith in the Moral Order of the World. By the late Baron BUNSEN. Translated from the German by SUSANNA WINKWORTH; with a Preface by Dean STANLEY. In Three Volumes. Vols. I. and II. 8vo. 30s.

HISTORY of EUROPEAN MORALS, from Augustus to Charlemagne. By W. E. H. LECKY, M.A. 2 vols. 8vo. price 23s.

HISTORY of the RISE and INFLUENCE of the SPIRIT of RATIONALISM in EUROPE. By W. E. H. LECKY, M.A. Third Edition, revised. 2 vols. 8vo. 25s.

The HISTORY of PHILOSOPHY, from Thales to Comte. By GEORGE HENRY LEWES. Third Edition. 2 vols. 8vo. 30s.

EGYPT'S PLACE in UNIVERSAL HISTORY; an Historical Investigation. By Baron BUNSEN, D.C.L. Translated by C. H. COTTRELL, M.A. With Additions by S. BIERCH, LL.D. 5 vols. 8vo. price £3 14s. 6d.

- MAUNDER'S HISTORICAL TREASURY**; comprising a General Introductory Outline of Universal History, and a series of Separate Histories. Latest Edition, revised and brought down to the Present Time by the Rev. G. W. COX, M.A. late Scholar of Trinity College, Oxford. Fcp. 10s. 6d.
- HISTORY of the NORMAN KINGS of ENGLAND.** Drawn from a New Collation of the Contemporary Chronicles, by THOMAS COBBE, of the Inner Temple, Barrister-at-Law. 1 vol. 8vo. [Nearly ready.]
- HISTORY of the CHRISTIAN CHURCH**, from the Ascension of Christ to the Conversion of Constantine. By E. BURTON, D.D. late Prof. of Divinity in the Univ. of Oxford. Eighth Edition. Fcp. 3s. 6d.
- SKETCH of the HISTORY of the CHURCH of ENGLAND to the Revolution of 1688.** By the Right Rev. T. V. SHORT, D.D. Lord Bishop of St. Asaph. Seventh Edition. Crown 8vo. 10s. 6d.
- HISTORY of the EARLY CHURCH**, from the First Preaching of the Gospel to the Council of Nicæa, A.D. 325. By ELIZABETH M. SEWELL, Author of 'Amy Herbert.' Fcp. 4s. 6d.
- The ENGLISH REFORMATION.** By 'F. C. MASSINGBERD, M.A. Chancellor of Lincoln and Rector of South Ormsby. Fourth Edition, revised. Fcp. 8vo. 7s. 6d.

Biography and Memoirs.

- DICTIONARY of GENERAL BIOGRAPHY**; containing Concise Memoirs and Notices of the most Eminent Persons of all Countries, from the Earliest Ages to the Present Time. Edited by W. L. E. CATES. 8vo. 21s.
- LIVES of the TUDOR PRINCESSES**, including Lady Jane Grey and her Sisters. By AGNES STRICKLAND, Author of 'Lives of the Queens of England.' Post 8vo. with Portrait, &c. 12s. 6d.
- MEMOIRS of BARON BUNSEN.** Drawn chiefly from Family Papers by his Widow, FRANCES BARONESS BUNSEN. Second Edition, abridged; with 2 Portraits and 4 Woodcuts. 2 vols. post 8vo. 21s.
- LIFE and CORRESPONDENCE of RICHARD WHATELY, D.D.** late Archbishop of Dublin. By E. JANE WHATELY. Popular Edition, with Additions and Omissions. Crown 8vo. with Portrait, 7s. 6d.
- LIFE of the DUKE of WELLINGTON.** By the Rev. G. R. GLEIG, M.A. Popular Edition, carefully revised; with copious Additions. Crown 8vo. with Portrait, 5s.
- HISTORY of MY RELIGIOUS OPINIONS.** By J. H. NEWMAN, D.D. Being the Substance of Apologia pro Vita Sua. Post 8vo. 6s.
- FATHER MATHEW: a Biography.** By JOHN FRANCIS MAGUIRE, M.P. for Cork. Popular Edition, with Portrait. Crown 8vo. 3s. 6d.
- THE LIFE of FRANZ SCHUBERT.** Translated from the German of K. VON HELLBORN, by A. D. COLERIDGE, M.A. late Fellow of King's College, Cambridge. With an APPENDIX by G. GROVE. 2 vols. post 8vo. with Portrait, 21s.

REMINISCENCES of FELIX MENDELSSOHN-BARTHOLDY; a Social and Artistic Biography. By ELISE POLKO. Translated from the German by Lady WALLACE. With additional Letters addressed to English Correspondents. Post 8vo. with Portrait and View, 10s. 6d.

FELIX MENDELSSOHN'S LETTERS from *Italy and Switzerland*, and *Letters from 1833 to 1847*, translated by Lady WALLACE. New Edition, with Portrait. 2 vols. crown 8vo. 5s. each.

FARADAY as a DISCOVERER. By JOHN TYNDALL, LL.D. F.R.S. Professor of Natural Philosophy in the Royal Institution of Great Britain. With Two Portraits. Crown 8vo. 6s.

MEMOIRS of SIR HENRY HAVELOCK, K.C.B. By JOHN CLARK MARSHMAN. Cabinet Edition, with Portrait. Crown 8vo. price 5s.

CAPTAIN COOK'S LIFE, VOYAGES, and DISCOVERIES. 18mo. Woodcuts, 2s. 6d.

LIFE of Sir JOHN RICHARDSON, C.B. sometime Inspector of Naval Hospitals and Fleets. By the Rev. JOHN MCILRAITH. Fcp. 8vo. with Portrait, 5s.

LIFE of PASTOR FLIEDNER, Founder of the Deaconesses' Institution at Kaiserswerth. Translated from the German by CATHERINE WINKWORTH. Fcp. 8vo. with Portrait, 3s. 6d.

VICISSITUDES of FAMILIES. By Sir J. BERNARD BURKE, C.B. Ulster King of Arms. New Edition, remodelled and enlarged. 2 vols. crown 8vo. 21s.

THE EARLS of GRANARD: a Memoir of the Noble Family of Forbes. Written by Admiral the Hon. JOHN FORBES, and edited by GEORGE ARTHUR HASTINGS, present Earl of Granard, K.P. 8vo. 10s.

GEORGE PETRIE, LL.D. M.R.I.A. &c. formerly President of the Royal Hibernian Academy; his Life and Labours in Art and Archæology. By WILLIAM STOKES, M.D. &c. Physician-in-Ordinary to the Queen in Ireland. 8vo. 12s. 6d.

ESSAYS in ECCLESIASTICAL BIOGRAPHY. By the Right Hon. Sir J. STEPHEN, LL.D. Cabinet Edition (being the Fifth). Crown 8vo. 7s. 6d.

ESSAYS on EDUCATIONAL REFORMERS: the Jesuits, Locke, J. J. Rousseau, Pestalozzi, Jacotot, &c. By the Rev. R. H. QUICK, M.A. Trin. Coll. Cantab. Post 8vo. 7s. 6d.

ESSAYS, BIOGRAPHICAL and CRITICAL. By A. L. MEISSNER, Ph.D. Professor of Modern Languages in Queen's College, Belfast, and in the Queen's University in Ireland. [Nearly ready.]

MAUNDER'S BIOGRAPHICAL TREASURY. Thirteenth Edition, reconstructed, thoroughly revised, and in great part rewritten; with about 1,000 additional Memoirs and Notices, by W. L. R. CATES. Fcp. 10s. 6d.

LETTERS and LIFE of FRANCIS BACON, including all his Occasional Works. Collected and edited, with a Commentary, by J. SPEDDING, Trin. Coll. Cantab. Vols. I. and II. 8vo. 24s. Vols. III. and IV. price 24s.

Criticism, Philosophy, Polity, &c.

THE INSTITUTES OF JUSTINIAN; with English Introduction, Translation, and Notes. By T. C. SANDARS, M.A. Barrister, late Fellow of Oriel Coll. Oxon. Fourth Edition. 8vo. 15s.

SOCRATES and the SOCRATIC SCHOOLS. Translated from the German of Dr. E. ZELLER, with the Author's approval, by the Rev. OSWALD J. REICHEL, B.C.L. and M.A. Crown 8vo. 8s. 6d.

THE ETHICS OF ARISTOTLE, illustrated with Essays and Notes. By Sir A. GRANT, Bart. M.A. LL.D. Second Edition, revised and completed. 2 vols. 8vo. price 28s.

ELEMENTS OF LOGIC. By R. WHATELY, D.D. late Archbishop of Dublin. Ninth Edition. 8vo. 10s. 6d. crown 8vo. 4s. 6d.

Elements of Rhetoric. By the same Author. Seventh Edition. 8vo, 10s. 6d. crown 8vo. 4s. 6d.

English Synonymes. By E. JANE WHATELY. Edited by Archbishop WHATELY. 5th Edition. Fcp. 3s.

BACON'S ESSAYS with ANNOTATIONS. By R. WHATELY, D.D. late Archbishop of Dublin. Sixth Edition. 8vo. 10s. 6d.

LORD BACON'S WORKS, collected and edited by R. L. ELLIS, M.A. J. SPEDDING, M.A. and D. D. HEATH. Vols. I. to V. *Philosophical Works*, 5 vols. 8vo. £4 6s. Vols. VI. and VII. *Literary and Professional Works*, 2 vols. £1 16s.

ON REPRESENTATIVE GOVERNMENT. By JOHN STUART MILL. Third Edition. 8vo. 9s. Crown 8vo. 2s.

ON LIBERTY. By JOHN STUART MILL. Fourth Edition. Post 8vo. 7s. 6d. Crown 8vo. 1s. 4d.

Principles of Political Economy. By the same Author. Sixth Edition. 2 vols. 8vo. 30s. Or in 1 vol. crown 8vo. 5s.

A System of Logic, Ratiocinative and Inductive. By the same Author. Seventh Edition. Two vols. 8vo. 25s.

ANALYSIS OF MR. MILL'S SYSTEM OF LOGIC. By W. STEBBING, M.A. Fellow of Worcester College, Oxford. Second Edition. 12mo. 3s. 6d.

UTILITARIANISM. By JOHN STUART MILL. Third Edition. 8vo. 5s.

Dissertations and Discussions, Political, Philosophical, and Historical. By the same Author. Second Edition, revised. 3 vols. 8vo. 36s.

Examination of Sir W. Hamilton's Philosophy, and of the Principal Philosophical Questions discussed in his Writings. By the same Author. Third Edition. 8vo. 16s.

AN OUTLINE OF THE NECESSARY LAWS OF THOUGHT: a Treatise on Pure and Applied Logic. By the Most Rev. WILLIAM, Lord Archbishop of York, D.D. F.R.S. Ninth Thousand. Crown 8vo. 5s. 6d.

The ELEMENTS of POLITICAL ECONOMY. By HENRY DUNNING MACLEOD, M.A. Barrister-at-Law. 8vo. 18s.

A Dictionary of Political Economy; Biographical, Bibliographical, Historical, and Practical. By the same Author. VOL. I. royal 8vo. 30s.

The ELECTION of REPRESENTATIVES, Parliamentary and Municipal; a Treatise. By THOMAS HARE, Barrister-at-Law. Third Edition, with Additions. Crown 8vo. 6s.

SPEECHES of the RIGHT HON. LORD MACAULAY, corrected by Himself. Library Edition, 8vo. 12s. People's Edition, crown 8vo. 3s. 6d.

LORD MACAULAY'S SPEECHES on PARLIAMENTARY REFORM in 1831 and 1832. 16mo. 1s.

INAUGURAL ADDRESS delivered to the University of St. Andrews. By JOHN STUART MILL. 8vo. 5s. People's Edition, crown 8vo. 1s.

A DICTIONARY of the ENGLISH LANGUAGE. By R. G. LATHAM, M.A. M.D. F.R.S. Founded on the Dictionary of Dr. SAMUEL JOHNSON, as edited by the Rev. H. J. TODD, with numerous Emendations and Additions. In Two Volumes. VOL. I. 4to. in Two Parts, price £3 10s. In course of publication, also, in 36 Parts, price 3s. 6d. each.

THESAURUS of ENGLISH WORDS and PHRASES, classified and arranged so as to facilitate the Expression of Ideas, and assist in Literary Composition. By P. M. ROBERT, M.D. New Edition. Crown 8vo. 10s. 6d.

LECTURES on the SCIENCE of LANGUAGE, delivered at the Royal Institution. By MAX MÜLLER, M.A. Fellow of All Souls College, Oxford. 2 vols. 8vo. FIRST SERIES, Fifth Edition, 12s. SECOND SERIES, Second Edition, 18s.

CHAPTERS on LANGUAGE. By FREDERIC W. FARRAR, F.R.S. late Fellow of Trin. Coll. Cambridge. Crown 8vo. 8s. 6d.

WORD-GOSSIP; a Series of Familiar Essays on Words and their Peculiarities. By the Rev. W. L. BLACKLEY, M.A. Fcp. 8vo. 5s.

A BOOK ABOUT WORDS. By G. F. GRAHAM, Author of 'English, or the Art of Composition,' 'English Synonymes,' 'English Grammar Practice,' 'English Style,' &c. Fcp. 8vo. [Nearly ready.]

The DEBATER; a Series of Complete Debates, Outlines of Debates, and Questions for Discussion. By F. ROWTON. Fcp. 6s.

MANUAL of ENGLISH LITERATURE, Historical and Critical. By THOMAS ARNOLD, M.A. Second Edition. Crown 8vo. price 7s. 6d.

SOUTHEY'S DOCTOR, complete in One Volume. Edited by the Rev. J. W. WARTER, B.D. Square crown 8vo. 12s. 6d.

HISTORICAL and CRITICAL COMMENTARY on the OLD TESTAMENT; with a New Translation. By M. M. KALISCH, Ph.D. VOL. I. *Genesis*, 8vo. 18s. or adapted for the General Reader, 12s. VOL. II. *Exodus*, 15s. or adapted for the General Reader, 12s. VOL. III. *Leviticus*, PART I. 15s. or adapted for the General Reader, 8s.

A Hebrew Grammar, with Exercises. By the same Author. PART I. *Outlines with Exercises*, 8vo. 12s. 6d. KEY, 5s. PART II. *Exceptional Forms and Constructions*, 12s. 6d.

- A LATIN-ENGLISH DICTIONARY.** By J. T. WHITE, D.D. of Corpus Christi College, and J. E. RIDDLE, M.A. of St. Edmund Hall, Oxford. 2 vols. 4to. pp. 2,128, price 42s. cloth.
- White's College Latin-English Dictionary** (Intermediate Size), abridged for the use of University Students from the Parent Work (as above). Medium 8vo. pp. 1,048, price 18s. cloth.
- White's Junior Student's College Latin-English and English-Latin Dictionary.** Square 12mo. pp. 1,058, price 12s.
- Separately { The ENGLISH-LATIN DICTIONARY, price 5s. 6d.
The LATIN-ENGLISH DICTIONARY, price 7s. 6d.
- An ENGLISH-GREEK LEXICON**, containing all the Greek Words used by Writers of good authority. By C. D. YONGE, B.A. New Edition. 4to. 21s.
- Mr. YONGE'S NEW LEXICON**, English and Greek, abridged from his larger work (as above). Revised Edition. Square 12mo. 8s. 6d.
- A GREEK-ENGLISH LEXICON.** Compiled by H. G. LIDDELL, D.D. Dean of Christ Church, and R. SCOTT, D.D. Master of Balliol. Fifth Edition. Crown 4to. 31s. 6d.
- A Lexicon, Greek and English**, abridged from LIDDELL and SCOTT'S *Greek-English Lexicon*. Twelfth Edition. Square 12mo. 7s. 6d.
- A SANSKRIT-ENGLISH DICTIONARY**, the Sanskrit words printed both in the original Devanagari and in Roman Letters. Compiled by T. BENFET, Prof. in the Univ. of Göttingen. 8vo. 52s. 6d.
- WALKER'S PRONOUNCING DICTIONARY** of the ENGLISH LANGUAGE. Thoroughly revised Editions, by B. H. SMART. 8vo. 12s. 16mo. 6s.
- A PRACTICAL DICTIONARY** of the FRENCH and ENGLISH LANGUAGES. By L. CONTANSEAU. Thirteenth Edition. Post 8vo. 10s. 6d.
- Contanseau's Pocket Dictionary**, French and English, abridged from the above by the Author. New Edition, revised. Square 18mo. 3s. 6d.
- NEW PRACTICAL DICTIONARY** of the GERMAN LANGUAGE; German-English and English-German. By the Rev. W. L. BLACKLEY, M.A. and Dr. CARL MARTIN FRIEDLÄNDER. Post 8vo. 7s. 6d.

Miscellaneous Works and Popular Metaphysics.

- The ESSAYS and CONTRIBUTIONS** of A. K. H. B., Author of 'The Recreations of a Country Parson.' Uniform Editions:—
- Recreations of a Country Parson.** FIRST and SECOND SERIES, crown 8vo. 3s. 6d. each.
- The Common-place Philosopher in Town and Country.** Crown 8vo. 3s. 6d.
- Leisure Hours in Town; Essays** Consolatory, Æsthetical, Moral, Social, and Domestic. Crown 8vo. 3s. 6d.

The Autumn Holidays of a Country Parson; Essays contributed to *Fraser's Magazine* and to *Good Words*. Crown 8vo. 3s. 6d.

The Graver Thoughts of a Country Parson. FIRST and SECOND SERIES, crown 8vo. 3s. 6d. each.

Critical Essays of a Country Parson. Selected from Essays contributed to *Fraser's Magazine*. Crown 8vo. 3s. 6d.

Sunday Afternoons at the Parish Church of a Scottish University City. Crown 8vo. 3s. 6d.

Lessons of Middle Age, with some Account of various Cities and Men. By A. K. H. B. Author of 'The Recreations of a Country Parson.' Crown 8vo. 3s. 6d.

Counsel and Comfort spoken from a City Pulpit. Crown 8vo. 3s. 6d.

Changed Aspects of Unchanged Truths: Memorials of St. Andrews Sundays. Crown 8vo. 3s. 6d.

SHORT STUDIES on GREAT SUBJECTS. By JAMES ANTHONY FROUDE, M.A. late Fellow of Exeter Coll. Oxford. Third Edition. 8vo. 12s.

LORD MACAULAY'S MISCELLANEOUS WRITINGS:—

LIBRARY EDITION. 2 vols. 8vo. Portrait, 21s.

PEOPLE'S EDITION. 1 vol. crown 8vo. 4s. 6d.

The REV. SYDNEY SMITH'S MISCELLANEOUS WORKS; including his Contributions to the *Edinburgh Review*. 2 vols. crown 8vo. 8s.

The Wit and Wisdom of the Rev. Sydney Smith. a Selection of the most memorable Passages in his Writings and Conversation. 16mo. 5s.

EPIGRAMS, Ancient and Modern; Humorous, Witty, Satirical, Moral, and Panegyrical. Edited by Rev. JOHN BOOTH, B.A. Cambridge. Second Edition, revised and enlarged. Fcp. 7s. 6d.

The PEDIGREE of the ENGLISH PEOPLE; an Argument, Historical and Scientific, on the *Ethnology* of the English. By THOMAS NICHOLAS, M.A. Ph.D. 8vo. 16s.

The ENGLISH and THEIR ORIGIN: a Prologue to authentic English History. By LUKE OWEN PIKE, M.A. Barrister-at-Law. 8vo. 9s.

ESSAYS selected from CONTRIBUTIONS to the *Edinburgh Review*. By HENRY ROGERS. Second Edition. 3 vols. fcp. 21s.

Reason and Faith, their Claims and Conflicts. By the same Author. New Edition, accompanied by several other Essays. Crown 8vo. 6s. 6d.

The Eclipse of Faith; or, a Visit to a Religious Sceptic. By the same Author. Twelfth Edition. Fcp. 5s.

Defence of the Eclipse of Faith, by its Author; a rejoinder to Dr. Newman's *Reply*. Third Edition. Fcp. 3s. 6d.

Selections from the Correspondence of R. E. H. Greyson. By the same Author. Third Edition. Crown 8vo. 7s. 6d.

CHIPS from a GERMAN WORKSHOP; being Essays on the Science of Religion, and on Mythology, Traditions, and Customs. By MAX MÜLLER, M.A. Fellow of All Souls College, Oxford. Second Edition, revised, with an Index. 2 vols. 8vo. 24s.

ANALYSIS of the PHENOMENA of the HUMAN MIND. By JAMES MILL. A New Edition, with Notes, Illustrative and Critical, by ALEXANDER BAIN, ANDREW FINDLATER, and GEORGE GROTE. Edited, with additional Notes, by JOHN STUART MILL. 2 vols. 8vo. price 28s.

AN INTRODUCTION to MENTAL PHILOSOPHY, on the Inductive Method. By J. D. MORELL, M.A. LL.D. 8vo. 12s.

Elements of Psychology, containing the Analysis of the Intellectual Powers. By the same Author. Post 8vo. 7s. 6d.

The SECRET of HEGEL: being the Hegelian System in Origin, Principle, Form, and Matter. By J. H. STIRLING. 2 vols. 8vo. 28s.

THE SENSES and the INTELLECT. By ALEXANDER BAIN, M.A. Professor of Logic in the University of Aberdeen. Third Edition. 8vo. 15s.

THE EMOTIONS and the WILL. By the same Author. Second Edition. 8vo. 15s.

On the STUDY of CHARACTER, including an Estimate of Phrenology. By the same Author. 8vo. 8s.

MENTAL and MORAL SCIENCE: a Compendium of Psychology and Ethics. By the same Author. Second Edition. Crown 8vo. 10s. 6d.

THE PHILOSOPHY of NECESSITY; or, Natural Law as applicable to Mental, Moral, and Social Science. By CHARLES BRAY. Second Edition. 8vo. 9s.

The Education of the Feelings and Affections. By the same Author. Third Edition. 8vo. 3s. 6d.

On Force, its Mental and Moral Correlates. By the same Author. 8vo. 5s.

THE FOLK-LORE of the NORTHERN COUNTIES of ENGLAND and the Borders. By WILLIAM HENDERSON. With an Appendix on Household Stories by the Rev. S. BARING-GOULD, M.A. Post 8vo. 9s. 6d.

Astronomy, Meteorology, Popular Geography, &c.

OUTLINES of ASTRONOMY. By Sir J. F. W. HERSCHEL, Bart. M.A. Ninth Edition, revised; with Plates and Woodcuts. 8vo. 18s.

SATURN and its SYSTEM. By RICHARD A. PROCTOR, B.A. late Scholar of St John's Coll. Camb. 8vo. with 14 Plates, 14s.

Handbook of the Stars. By the same Author. With 3 Maps. Square fcp. 5s.

CELESTIAL OBJECTS for COMMON TELESCOPES. By the Rev. T. W. WEBB, M.A. F.R.A.S. Second Edition, revised, with a large Map of the Moon, and several Woodcuts. 16mo. 7s. 6d.

NAVIGATION and NAUTICAL ASTRONOMY (Practical, Theoretical, Scientific) for the use of Students and Practical Men. By J. MERRIFIELD, F.R.A.S. and H. EVERS. 8vo. 14s.

- DOVE'S LAW of STORMS**, considered in connexion with the Ordinary Movements of the Atmosphere. Translated by R. H. SCOTT, M.A. T.C.D. 8vo. 10s. 6d.
- PHYSICAL GEOGRAPHY for SCHOOLS and GENERAL READERS.** By M. F. MAURY, LL.D. Fcp. with 2 Charts, 2s. 6d.
- A TREATISE on the ACTION of VIS INERTIE in the OCEAN;** with Remarks on the Abstract Nature of the Forces of Vis Inertiæ and Gravitation, and a New Theory of the Tides. By WILLIAM LEIGHTON JORDAN, F.R.G.S. With 12 Charts and Diagrams. 8vo. 14s.
- MC'ULLOCH'S DICTIONARY**, Geographical, Statistical, and Historical, of the various Countries, Places, and Principal Natural Objects in the World. New Edition, with the Statistical Information brought up to the latest returns by F. MARTIN. 4 vols. 8vo. with coloured Maps, £4 4s.
- A GENERAL DICTIONARY of GEOGRAPHY**, Descriptive, Physical, Statistical, and Historical: forming a complete Gazetteer of the World. By A. KEITH JOHNSTON, LL.D. F.R.G.S. Revised Edition. 8vo. 31s. 6d.
- A MANUAL of GEOGRAPHY**, Physical, Industrial, and Political. By W. HUGHES, F.R.G.S. With 6 Maps. Fcp. 7s. 6d.
- THE STATES of the RIVER PLATE:** their Industries and Commerce. By WILFRID LATHAM, Buenos Ayres. Second Edition, revised. 8vo. 12s.
- MAUNDER'S TREASURY of GEOGRAPHY**, Physical, Historical, Descriptive, and Political. Edited by W. HUGHES, F.R.G.S. With 7 Maps and 16 Plates. Fcp. 10s. 6d.

Natural History and Popular Science.

- ELEMENTARY TREATISE on PHYSICS**, Experimental and Applied. Translated and edited from GANOT's *Éléments de Physique* (with the Author's sanction) by E. ATKINSON, Ph. D. F.C.S. New Edition, revised and enlarged; with a Coloured Plate and 620 Woodcuts. Post 8vo. 15s.
- THE ELEMENTS of PHYSICS or NATURAL PHILOSOPHY.** By NEIL ARNOTT, M.D. F.R.S. Physician Extraordinary to the Queen. Sixth Edition, rewritten and completed. Two Parts, 8vo. 21s.
- SOUND:** a Course of Eight Lectures delivered at the Royal Institution of Great Britain. By JOHN TYNDALL, LL.D. F.R.S. Crown 8vo. with Portrait of *M. Chladni* and 169 Woodcuts, price 9s.
- HEAT CONSIDERED as a MODE of MOTION.** By Professor JOHN TYNDALL, LL.D. F.R.S. Third Edition. Crown 8vo. with Woodcuts, 10s. 6d.
- LIGHT:** Its Influence on Life and Health. By FORBES WINSLOW, M.D. D.C.L. Oxon. (Hon.). Fcp. 8vo. 6s.
- An ESSAY on DEW**, and several Appearances connected with it. By W. C. WELLS. Edited, with Annotations, by L. P. CASELLA, F.R.A.S. and an Appendix by E. STRACHAN, F.M.S. 8vo. 5s.
- A TREATISE on ELECTRICITY**, in Theory and Practice. By A. DE LA RIVE, Prof. in the Academy of Geneva. Translated by C. V. WALKER, F.R.S. 3 vols. 8vo. with Woodcuts, £3 13s.

- THE CORRELATION of PHYSICAL FORCES.** By W. R. GROVE, Q.O. V.P.R.S. Fifth Edition, revised, and followed by a Discourse on Continuity. 8vo. 10s. 6d. The *Discourse on Continuity*, separately, 2s. 6d.
- MANUAL of GEOLOGY.** By S. HAUGHTON, M.D. F.R.S. Revised Edition, with 66 Woodcuts. Fcp. 7s. 6d.
- A GUIDE to GEOLOGY.** By J. PHILLIPS, M.A. Professor of Geology in the University of Oxford. Fifth Edition, with Plates. Fcp. 4s.
- THE STUDENT'S MANUAL of ZOOLOGY and COMPARATIVE PHYSIOLOGY.** By J. BURNBY YEO, M.B. Resident Medical Tutor and Lecturer on Animal Physiology in King's College, London. [*Nearly ready.*]
- VAN DER HOEVEN'S HANDBOOK of ZOOLOGY.** Translated from the Second Dutch Edition by the Rev. W. CLARK, M.D. F.R.S. 2 vols. 8vo. with 24 Plates of Figures, 60s.
- Professor OWEN'S LECTURES on the COMPARATIVE ANATOMY and Physiology of the Invertebrate Animals.** Second Edition, with 235 Woodcuts. 8vo. 21s.
- THE COMPARATIVE ANATOMY and PHYSIOLOGY of the VERTEBRATE Animals.** By RICHARD OWEN, F.R.S. D.C.L. With 1,472 Woodcuts. 3 vols. 8vo. £3 13s. 6d.
- THE FIRST MAN and HIS PLACE in CREATION,** considered on the Principles of Common Sense from a Christian Point of View. By GEORGE MOORE, M.D. Post 8vo. 8s. 6d.
- THE PRIMITIVE INHABITANTS of SCANDINAVIA:** containing a Description of the Implements, Dwellings, Tombs, and Mode of Living of the Savages in the North of Europe during the Stone Age. By SVEN NILSSON. Translated from the Third Edition; with an Introduction by Sir J. LUBBOCK. With 16 Plates of Figures and 3 Woodcuts. 8vo. 18s.
- BIBLE ANIMALS;** being an Account of the various Birds, Beasts, Fishes, and other Animals mentioned in the Holy Scriptures. By the Rev. J. G. WOOD, M.A. F.L.S. Copiously illustrated with Original Designs, made under the Author's superintendence and engraved on Wood. In course of publication monthly, to be completed in 20 Parts, price 1s. each, forming One Volume, uniform with 'Homes without Hands.'
- HOMES WITHOUT HANDS:** a Description of the Habitations of Animals, classed according to their Principle of Construction. By Rev. J. G. WOOD, M.A. F.L.S. With about 140 Vignettes on Wood (20 full size of page). New Edition. 8vo. 21s.
- MANUAL of CORALS and SEA JELLIES.** By J. R. GREENE, B.A. Edited by JOSEPH A. GALBRAITH, M.A. and SAMUEL HAUGHTON, M.D. Fcp. with 39 Woodcuts, 5s.
- Manual of Sponges and Animalculæ;** with a General Introduction on the Principles of Zoology. By the same Author and Editors. Fcp. with 16 Woodcuts, 2s.
- Manual of the Metalloids.** By J. APJOHN, M.D. F.R.S. and the same Editors. Revised Edition. Fcp. with 38 Woodcuts, 7s. 6d.
- A FAMILIAR HISTORY of BIRDS.** By E. STANLEY, D.D. F.R.S. late Lord Bishop of Norwich. Seventh Edition, with Woodcuts. Fcp. 3s. 6d.

- The HARMONIES of NATURE and UNITY of CREATION.** By Dr. GEORGE HARTWIG. 8vo. with numerous Illustrations, 18s.
- The Sea and its Living Wonders.** By the same Author. Third (English) Edition. 8vo. with many Illustrations, 21s.
- The Tropical World.** By the same Author. With 8 Chromoxylographs and 172 Woodcuts. 8vo. 21s.
- The POLAR WORLD;** a Popular Description of Man and Nature in the Arctic and Antarctic Regions of the Globe. By Dr. GEORGE HARTWIG. With 8 Chromoxylographs, 3 Maps, and 85 Woodcuts. 8vo. 21s.
- CEYLON.** By Sir J. EMERSON TENNENT, K.C.S. LL.D. Fifth Edition; with Maps, &c. and 90 Wood Engravings. 2 vols. 8vo. £2 10s.
- KIRBY and SPENCE'S INTRODUCTION to ENTOMOLOGY,** or Elements of the Natural History of Insects. 7th Edition. Crown 8vo. 5s.
- MAUNDER'S TREASURY of NATURAL HISTORY,** or Popular Dictionary of Zoology. Revised and corrected by T. S. COBBOLD, M.D. Fcp. with 900 Woodcuts, 10s. 6d.
- The TREASURY of BOTANY,** or Popular Dictionary of the Vegetable Kingdom; including a Glossary of Botanical Terms. Edited by J. LINDLEY, F.R.S. and T. MOORE, F.L.S. assisted by eminent Contributors. Pp. 1,274, with 274 Woodcuts and 20 Steel Plates. 2 Parts. fcp. 20s.
- The ELEMENTS of BOTANY for FAMILIES and SCHOOLS.** Tenth Edition, revised by THOMAS MOORE, F.L.S. Fcp. with 154 Woodcuts, 2s. 6d.
- The ROSE AMATEUR'S GUIDE.** By THOMAS RIVERS. Twelfth Edition. Fcp. 4s.
- The BRITISH FLORA;** comprising the Phænogamous or Flowering Plants and the Ferns. By Sir W. J. HOOKER, K.H. and G. A. WALKER-ARNOTT, LL.D. 12mo. with 12 Plates, 14s. or coloured, 21s.
- LOUDON'S ENCYCLOPÆDIA of PLANTS;** comprising the Specific Character, Description, Culture, History, &c. of all the Plants found in Great Britain. With upwards of 12,000 Woodcuts. 8vo. 42s.
- MAUNDER'S SCIENTIFIC and LITERARY TREASURY.** New Edition, thoroughly revised and in great part re-written, with above 1,000 new Articles, by J. Y. JOHNSON, Corr. M.Z.S. Fcp. 10s. 6d.
- A DICTIONARY of SCIENCE, LITERATURE, and ART.** Fourth Edition, re-edited by W. T. BRANDE (the Author), and GEORGE W. COX, M.A. assisted by contributors of eminent Scientific and Literary Acquirements. 3 vols. medium 8vo. price 63s. cloth.
- The QUARTERLY JOURNAL of SCIENCE.** Edited by JAMES SAMUELSON and WILLIAM CROOKES, F.R.S. Published quarterly in January, April, July, and October. 8vo. with Illustrations, price 5s. each Number.

*Chemistry, Medicine, Surgery, and the
Allied Sciences.*

- A DICTIONARY of CHEMISTRY and the Allied Branches of other Sciences.** By HENRY WATTS, F.R.S. assisted by eminent Contributors. Complete in 5 vols. medium 8vo. £7 3s.

ELEMENTS of CHEMISTRY, Theoretical and Practical. By WILLIAM A. MILLER, M.D. LL.D. F.R.S. F.G.S. Prof. of Chemistry, King's Coll. London. 3 vols. 8vo. £3. PART I. CHEMICAL PHYSICS, 15s. PART II. INORGANIC CHEMISTRY, 21s. PART III. ORGANIC CHEMISTRY, 24s.

A MANUAL of CHEMISTRY, Descriptive and Theoretical. By WILLIAM ODLING, M.B. F.R.S. PART I. 8vo. 9s. PART II. *just ready*.

A Course of Practical Chemistry, for the use of Medical Students. By the same Author. New Edition, with 70 Woodcuts. Crown 8vo. 7s. 6d.

Lectures on Animal Chemistry, delivered at the Royal College of Physicians in 1865. By the same Author. Crown 8vo. 4s. 6d.

HANDBOOK of CHEMICAL ANALYSIS, adapted to the UNITARY System of Notation. By F. T. CONINGTON, M.A. F.C.S. Post 8vo. 7s. 6d. —CONINGTON's *Tables of Qualitative Analysis*, price 2s. 6d.

The DIAGNOSIS, PATHOLOGY, and TREATMENT of DISEASES of Women; including the Diagnosis of Pregnancy. By GRAILY HEWITT, M.D. Second Edition, enlarged; with 116 Woodcut Illustrations. 8vo. 24s.

LECTURES on the DISEASES of INFANCY and CHILDHOOD. By CHARLES WEST, M.D. &c. Fifth Edition, revised and enlarged. 8vo. 16s.

A SYSTEM of SURGERY, Theoretical and Practical. In Treatises by Various Authors. Edited by T. HOLMES, M.A. &c. Surgeon and Lecturer on Surgery at St. George's Hospital, and Surgeon-in-Chief to the Metropolitan Police. 4 vols. 8vo. £413s.

The SURGICAL TREATMENT of CHILDREN'S DISEASES. By T. HOLMES, M.A. &c. late Surgeon to the Hospital for Sick Children. Second Edition, with 9 Plates and 112 Woodcuts. 8vo. 21s.

LECTURES on the PRINCIPLES and PRACTICE of PHYSIC. By Sir THOMAS WATSON, Bart. M.D. New Edition in preparation.

LECTURES on SURGICAL PATHOLOGY. By J. PAGET, F.R.S. Edited by W. TURNER, M.B. New Edition in preparation.

On CHRONIC BRONCHITIS, especially as connected with GOUT, EMPHYSEMA, and DISEASES of the HEART. By E. HEADLAM GREENHOW, M.D. F.R.C.P. &c. 8vo. 7s. 6d.

A TREATISE on the CONTINUED FEVERS of GREAT BRITAIN. By C. MURCHISON, M.D. New Edition in preparation.

CLINICAL LECTURES on DISEASES of the LIVER, JAUNDICE, and ABDOMINAL DROPSY. By CHARLES MURCHISON, M.D. Post 8vo. with 25 Woodcuts, 10s. 6d.

ANATOMY, DESCRIPTIVE and SURGICAL. By HENRY GRAY, F.R.S. With 410 Wood Engravings from Dissections. New Edition, by T. HOLMES, M.A. Cantab. Royal 8vo. 28s.

The THEORY of OCULAR DEFECTS and of SPECTACLES. Translated from the German of Dr. H. SCHEFFLER by R. B. CARTER, F.R.C.S. With Prefatory Notes and a Chapter of Practical Instructions. Post 8vo. 7s. 6d.

OUTLINES of PHYSIOLOGY, Human and Comparative. By JOHN MARSHALL, F.R.C.S. Surgeon to the University College Hospital. 2 vols. crown 8vo. with 122 Woodcuts, 32s.

ESSAYS on PHYSIOLOGICAL SUBJECTS. By GILBERT W. CHILD, M.D. F.R.S. F.C.S. of Exeter College, Oxford. 8vo. 5s.

PHYSIOLOGICAL ANATOMY and PHYSIOLOGY of MAN. By the late R. B. TODD, M.D. F.R.S. and W. BOWMAN, F.R.S. of King's College. With numerous Illustrations. VOL. II. 8vo. 25s.

VOL. I. New Edition by Dr. LIONEL S. BEALE, F.R.S. in course of publication; PART I. with 8 Plates, 7s. 6d.

COPLAND'S DICTIONARY of PRACTICAL MEDICINE, abridged from the larger work and throughout brought down to the present State of Medical Science. 8vo. 38s.

The WORKS of SIR B. C. BRODIE, Bart. collected and arranged by CHARLES HAWKINS, F.R.C.S.E. 3 vols. 8vo. with Medallion and Facsimile, 48s.

On ANILINE and its DERIVATIVES: a Treatise on the Manufacture of Aniline and Aniline Colours. By M. REIMANN, Ph.D. L.A.M. To which is added the Report on the Colouring Matters derived from Coal Tar shewn at the French Exhibition of 1867. Edited by WILLIAM CROOKES, F.R.S. With 5 Woodcuts. 8vo. 10s. 6d.

A MANUAL of MATERIA MEDICA and THERAPEUTICS, abridged from Dr. PEREIRA's *Elements* by F. J. FARRE, M.D. assisted by R. BENTLEY, M.R.C.S. and by R. WARRINGTON, F.R.S. 8vo. with 90 Woodcuts, 21s.

THOMSON'S CONSPICUUS of the BRITISH PHARMACOPŒIA. 25th Edition, corrected by E. LLOYD BIRKETT, M.D. 18mo. price 6s.

MANUAL of the DOMESTIC PRACTICE of MEDICINE. By W. B. KESTEVEN, F.R.C.S.E. Third Edition, revised, with Additions. Fcp. 5s.

GYMNASTS and GYMNASTICS. By JOHN H. HOWARD, late Professor of Gymnastics, Comm. Coll. Rippondon. Second Edition, revised and enlarged, with 135 Woodcuts. Crown 8vo. 10s. 6d.

The Fine Arts, and Illustrated Editions.

MATERIALS for a HISTORY of OIL PAINTING. By Sir CHARLES LOCKE EASTLAKE, sometime President of the Royal Academy. VOL. II. 8vo. 14s.

HALF-HOUR LECTURES on the HISTORY and PRACTICE of the Fine and Ornamental Arts. By WILLIAM B. SCOTT. New Edition, revised by the Author; with 50 Woodcuts. Crown 8vo. 8s. 6d.

LECTURES on the HISTORY of MODERN MUSIC, delivered at the Royal Institution. By JOHN HULLAH. FIRST COURSE, with Chronological Tables, post 8vo. 6s. 6d. SECOND COURSE, on the Transition Period, with 40 Specimens, 8vo. 16s.

SIX LECTURES on HARMONY, delivered at the Royal Institution of Great Britain in the Year 1867. By G. A. MACFARREN. With numerous engraved Musical Examples and Specimens. 8vo. 10s. 6d.

The CHORALE BOOK for ENGLAND: the Hymns translated by Miss C. WINKWORTH; the tunes arranged by Prof. W. S. BENNETT and OTTO GOLDSCHMIDT. Fcp. 4to. 12s. 6d.

Congregational Edition. Fcp. 2s.

SACRED MUSIC for FAMILY USE; a Selection of Pieces for One, Two, or more Voices, from the best Composers, Foreign and English. Edited by JOHN HULLAH. 1 vol. music folio, price 21s.

The NEW TESTAMENT, illustrated with Wood Engravings after the Early Masters, chiefly of the Italian School. Crown 4to. 68s. cloth, gilt top; or 25 5s. elegantly bound in morocco.

LYRA GERMANICA; the Christian Year. Translated by CATHERINE WINKWORTH; with 125 Illustrations on Wood drawn by J. LEIGHTON, F.S.A. 4to. 21s.

LYRA GERMANICA; the Christian Life. Translated by CATHERINE WINKWORTH; with about 200 Woodcut Illustrations by J. LEIGHTON, F.S.A. and other Artists. 4to. 21s.

THE LIFE of MAN SYMBOLISED by the MONTHS of the YEAR. Text selected by R. PIGOT; Illustrations on Wood from Original Designs by J. LEIGHTON, F.S.A. 4to. 42s.

CATS' and FARLIE'S MORAL EMBLEMS; with Aphorisms, Adages, and Proverbs of all Nations. 121 Illustrations on Wood by J. LEIGHTON, F.S.A. Text selected by R. PIGOT. Imperial 8vo. 31s. 6d.

SHAKESPEARE'S MIDSUMMER NIGHT'S DREAM, illustrated with 24 Silhouettes or Shadow-Pictures by P. KONEWKA, engraved on Wood by A. VOGEL. Folio, 31s. 6d.

SHAKESPEARE'S SENTIMENTS and SIMILES, printed in Black and Gold, and Illuminated in the Missal Style by HENRY NOEL HUMPHREYS. Square post 8vo. 21s.

SACRED and LEGENDARY ART. By Mrs. JAMESON.

Legends of the Saints and Martyrs. Fifth Edition, with 19 Etchings and 187 Woodcuts. 2 vols. square crown 8vo. 31s. 6d.

Legends of the Monastic Orders. Third Edition, with 11 Etchings and 88 Woodcuts. 1 vol. square crown 8vo. 21s.

Legends of the Madonna. Third Edition, with 27 Etchings and 165 Woodcuts. 1 vol. square crown 8vo. 21s.

The History of Our Lord, with that of his Types and Precursors. Completed by Lady EASTLAKE. Revised Edition, with 31 Etchings and 281 Woodcuts. 2 vols. square crown 8vo. 42s.

Arts, Manufactures, &c.

DRAWING from NATURE. By GEORGE BARNARD, Professor of Drawing at Rugby School. With 18 Lithographic Plates, and 108 Wood Engravings. Imperial 8vo. price 25s. Or in Three Parts, royal 8vo. 7s. 6d. each.

GWILT'S ENCYCLOPEDIA of ARCHITECTURE, with above 1,100 Engravings on Wood. Fifth Edition, revised and enlarged by WYATT PAPWORTH. Additionally illustrated with nearly 400 Wood Engravings by O. Jewitt, and more than 100 other new Woodcuts. 8vo. 52s. 6d.

ITALIAN SCULPTORS; being a History of Sculpture in Northern, Southern, and Eastern Italy. By C. C. PERKINS. With 30 Etchings and 13 Wood Engravings. Imperial 8vo. 42s.

TUSCAN SCULPTORS, their Lives, Works, and Times. With 45 Etchings and 28 Woodcuts from Original Drawings and Photographs. By the same Author. 2 vols. imperial 8vo. 63s.

ORIGINAL DESIGNS for WOOD-CARVING, with **PRACTICAL INSTRUCTIONS** in the Art. By A. F. B. With 20 Plates of Illustrations engraved on Wood. 4to. 18s.

HINTS on HOUSEHOLD TASTE in FURNITURE, UPHOLSTERY, and other Details. By CHARLES L. EASTLAKE, Architect. With about 90 Illustrations. Square crown 8vo. 13s.

The ENGINEER'S HANDBOOK; explaining the Principles which should guide the Young Engineer in the Construction of Machinery. By C. S. LOWENDES. Post 8vo. 5s.

The ELEMENTS of MECHANISM. By T. M. GOODEVE, M.A. Professor of Mechanics at the E. M. Acad. Woolwich. Second Edition, with 217 Woodcuts. Post 8vo. 6s. 6d.

LATHES and TURNING, Simple, Mechanical, and **ORNAMENTAL**. By W. HENRY NORTHCOTT. With about 240 Illustrations on Steel and Wood. 8vo. 18s.

URE'S DICTIONARY of ARTS, MANUFACTURES, and MINES. Sixth Edition, chiefly rewritten and greatly enlarged by ROBERT HUNT, F.R.S. assisted by numerous Contributors eminent in Science and the Arts, and familiar with Manufactures. With above 2,000 Woodcuts. 3 vols. medium 8vo. price £4 14s. 6d.

HANDBOOK of PRACTICAL TELEGRAPHY, published with the sanction of the Chairman and Directors of the Electric and International Telegraph Company, and adopted by the Department of Telegraphs for India. By E. S. CULLEY. Third Edition. 8vo. 12s. 6d.

ENCYCLOPÆDIA of CIVIL ENGINEERING, Historical, Theoretical, and Practical. By E. CRESEY, C.E. With above 3,000 Woodcuts. 8vo. 42s.

TREATISE on MILLS and MILLWORK. By W. FAIRBAIRN, C.E. Second Edition, with 18 Plates and 322 Woodcuts. 2 vols. 8vo. 32s.

Useful Information for Engineers. By the same Author. **FIRST, SECOND, and THIRD SERIES**, with many Plates and Woodcuts. 3 vols. crown 8vo. 10s. 6d. each.

The Application of Cast and Wrought Iron to Building Purposes. By the same Author. Third Edition, with 6 Plates and 118 Woodcuts. 8vo. 16s.

IRON SHIP BUILDING, its History and Progress, as comprised in a Series of Experimental Researches. By the same Author. With 4 Plates and 130 Woodcuts. 8vo. 18s.

A TREATISE on the STEAM ENGINE, in its various Applications to Mines, Mills, Steam Navigation, Railways and Agriculture. By J. BOURNE, C.E. Eighth Edition; with Portrait, 37 Plates, and 546 Woodcuts. 4to. 42s.

Catechism of the Steam Engine, in its various Applications to Mines, Mills, Steam Navigation, Railways, and Agriculture. By the same Author. With 89 Woodcuts. Fcp. 6s.

Handbook of the Steam Engine. By the same Author, forming a **KEY** to the Catechism of the Steam Engine, with 67 Woodcuts. Fcp. 9s.

A TREATISE on the SCREW PROPELLER, SCREW VESSELS, and Screw Engines, as adapted for purposes of Peace and War; with Notices of other Methods of Propulsion, Tables of the Dimensions and Performance of Screw Steamers, and detailed Specifications of Ships and Engines. By J. BOURNE, C.E. Third Edition, with 54 Plates and 287 Woodcuts. 4to. 63s.

EXAMPLES of MODERN STEAM, AIR, and GAS ENGINES of the most Approved Types, as employed for Pumping, for Driving Machinery, for Locomotion, and for Agriculture, minutely and practically described. Illustrated by Working Drawings, and embodying a Critical Account of all Projects of Recent Improvement in Furnaces, Boilers, and Engines. By the same Author. In course of publication monthly, to be completed in 24 Parts, price 2s. 6d. each, forming One volume 4to. with about 50 Plates and 400 Woodcuts.

A HISTORY of the MACHINE-WROUGHT HOSIERY and LACE Manufactures. By WILLIAM FELKIN, F.L.S. F.R.S. Royal 8vo. 21s.

PRACTICAL TREATISE on METALLURGY, adapted from the last German Edition of Professor KERL's *Metallurgy* by W. CROOKES, F.R.S. &c. and E. RÖHRIG, Ph.D. M.E. Vol. I. comprising *Lead, Silver, Zinc, Cadmium, Tin, Mercury, Bismuth, Antimony, Nickel, Arsenic, Gold, Platinum, and Sulphur*. 8vo. with 207 Woodcuts, 31s. 6d.

MITCHELL'S MANUAL of PRACTICAL ASSAYING. Third Edition, for the most part re-written, with all the recent Discoveries incorporated, by W. CROOKES, F.R.S. With 188 Woodcuts. 8vo. 28s.

The ART of PERFUMERY; the History and Theory of Odours, and the Methods of Extracting the Aromas of Plants. By Dr. PIESSE, F.R.S. Third Edition, with 53 Woodcuts. Crown 8vo. 10s. 6d.

Chemical, Natural, and Physical Magic, for Juveniles during the Holidays. By the same Author. Third Edition, with 38 Woodcuts. Fcp. 6s.

LOUDON'S ENCYCLOPÆDIA of AGRICULTURE: comprising the Laying-out, Improvement, and Management of Landed Property, and the Cultivation and Economy of the Productions of Agriculture. With 1,100 Woodcuts. 8vo. 31s. 6d.

Loudon's Encyclopædia of Gardening: comprising the Theory and Practice of Horticulture, Floriculture, Arboriculture, and Landscape Gardening. With 1,000 Woodcuts. 8vo. 31s. 6d.

BAYLDON'S ART of VALUING RENTS and TILLAGES, and Claims of Tenants upon Quitting Farms, both at Michaelmas and Lady-Day. Eighth Edition, revised by J. C. MORTON. 8vo. 10s. 6d.

Religious and Moral Works.

An EXPOSITION of the 39 ARTICLES, Historical and Doctrinal. By E. HAROLD BROWNE, D.D. Lord Bishop of Ely. Seventh Edit. 8vo. 16s.

ARCHBISHOP LEIGHTON'S SERMONS and CHARGES. With Additions and Corrections from MSS. and with Historical and other Illustrative Notes by WILLIAM WEST, Incumbent of S. Columba's, Nairn. 8vo. 15s.

The ACTS of the APOSTLES; with a Commentary, and Practical and Devotional Suggestions for Readers and Students of the English Bible. By the Rev. F. C. COOK, M.A. Canon of Exeter, &c. New Edition. 8vo. 12s. 6d.

- The LIFE and EPISTLES of ST. PAUL.** By W. J. CONYBEARE, M.A. late Fellow of Trin. Coll. Cantab. and the Very Rev. J. S. HOWSON, D.D. Dean of Chester.
- LIBRARY EDITION, with all the Original Illustrations, Maps, Landscapes on Steel, Woodcuts, &c. 2 vols. 4to. 48s.
- INTERMEDIATE EDITION, with a Selection of Maps, Plates, and Woodcuts. 2 vols. square crown 8vo. 31s. 6d.
- PEOPLE'S EDITION, revised and condensed, with 46 Illustrations and Maps. 2 vols. crown 8vo. 12s.
- The VOYAGE and SHIPWRECK of ST. PAUL;** with Dissertations on the Life and Writings of St. Luke and the Ships and Navigation of the Ancients. By JAMES SMITH, F.R.S. Third Edition. Crown 8vo. 10s. 6d.
- The NATIONAL CHURCH; HISTORY and PRINCIPLES of the CHURCH POLITY of ENGLAND.** By D. MOUNTFIELD, M.A. Rector of Newport, Salop. Crown 8vo. 4s.
- EVIDENCE of the TRUTH of the CHRISTIAN RELIGION** derived from the Literal Fulfilment of Prophecy. By ALEXANDER KEITH, D.D. 37th Edition, with numerous Plates, in square 8vo. 12s. 6d.; also the 39th Edition, in post 8vo. with 5 Plates, 6s.
- The HISTORY and DESTINY of the WORLD and of the CHURCH.** according to Scripture. By the same Author. Square 8vo. with 40 Illustrations, 10s.
- A CRITICAL and GRAMMATICAL COMMENTARY on ST. PAUL'S Epistles.** By C. J. ELLICOTT, D.D. Lord Bishop of Gloucester & Bristol. 8vo. Galatians, Fourth Edition, 8s. 6d.
- Ephesians, Fourth Edition, 8s. 6d.
- Pastoral Epistles, Fourth Edition, 10s. 6d.
- Philippians, Colossians, and Philemon, Third Edition, 10s. 6d.
- Thessalonians, Third Edition, 7s. 6d.
- Historical Lectures on the Life of our Lord Jesus Christ:** being the Hulsean Lectures for 1859. By the same Author. Fourth Edition. 8vo. price 10s. 6d.
- An INTRODUCTION to the STUDY of the NEW TESTAMENT,** Critical, Exegetical, and Theological. By the Rev. S. DAVIDSON, D.D. LL.D. 2 vols. 8vo. 30s.
- Rev. T. H. HORNE'S INTRODUCTION to the CRITICAL STUDY** and Knowledge of the Holy Scriptures. Twelfth Edition, as last revised throughout and brought up to the existing state of Biblical Knowledge under careful Editorial revision. With 4 Maps and 23 Woodcuts and Facsimiles. 4 vols. 8vo. 42s.
- Rev. T. H. Horne's Compendious Introduction to the Study of the Bible,** being an Analysis of the larger work by the same Author. Re-edited by the Rev. JOHN AYRE, M.A. With Maps, &c. Post 8vo. 6s.
- EWALD'S HISTORY of ISRAEL to the DEATH of MOSES.** Translated from the German. Edited, with a Preface and an Appendix, by RUSSELL MARTINEAU, M.A. Prof. of Hebrew in Manchester New Coll. London. Second Edition, continued to the commencement of the Monarchy. 2 vols. 8vo. 24s. VOL. II. comprising *Joshua* and *Judges*, for Purchasers of the First Edition, 9s.

The TREASURY of BIBLE KNOWLEDGE; being a Dictionary of the Books, Persons, Places, Events, and other matters of which mention is made in Holy Scripture. By Rev. J. AYRE, M.A. With Maps, 16 Plates, and numerous Woodcuts. Fcp. 10s. 6d.

The GREEK TESTAMENT; with Notes, Grammatical and Exegetical. By the Rev. W. WEBSTER, M.A. and the Rev. W. F. WILKINSON, M.A. 2 vols. 8vo. £2 4s.

VOL. I. the Gospels and Acts, 20s.

VOL. II. the Epistles and Apocalypse, 24s.

The CHURCHMAN'S DAILY REMEMBRANCER of DOCTRINE and DUTY: consisting of Meditations taken from the Writings of Standard Divines from the Early Days of Christianity to the Present Time; with a Preface by W. R. FREMANTLE, M.A. New Edition. Fcp. 8vo. 6s.

EVERY-DAY SCRIPTURE DIFFICULTIES explained and illustrated. By J. E. PRESCOTT, M.A. VOL. I. *Matthew and Mark*; VOL. II. *Luke and John*. 2 vols. 8vo. 9s. each.

The PENTATEUCH and BOOK of JOSHUA CRITICALLY EXAMINED. By the Right Rev. J. W. COLENSO, D.D. Lord Bishop of Natal. People's Edition, in 1 vol. crown 8vo. 6s. or in 5 Parts, 1s. each.

The CHURCH and the WORLD: Three Series of Essays on Questions of the Day. By Various Writers. Edited by the Rev. ORBY SHIPLEY, M.A. FIRST SERIES, Third Edition, 15s. SECOND SERIES, Second Edition, 15s. THIRD SERIES, 1868, price 15s. 3 vols. 8vo. 45s.

The FORMATION of CHRISTENDOM. By T. W. ALLIEN. PARTS I. and II. 8vo. price 12s. each Part.

ENGLAND and CHRISTENDOM. By ARCHBISHOP MANNING, D.D. Post 8vo. price 10s. 6d.

CHRISTENDOM'S DIVISIONS, PART I., a Philosophical Sketch of the Divisions of the Christian Family in East and West. By EDMUND S. FFOULKES. Post 8vo. price 7s. 6d.

Christendom's Divisions, PART II. Greeks and Latins, being a History of their Dissensions and Overtures for Peace down to the Reformation. By the same Author. Post 8vo. 15s.

The WOMAN BLESSED by ALL GENERATIONS; or, Mary the Object of Veneration, Confidence, and Imitation to all Christians. By the Rev. R. MELIA, D.D. P.S.M. With 78 Illustrations. 8vo. 15s.

LIFE of the BLESSED VIRGIN; the FEMALL GLORY. By ANTHONY STAFFORD. Together with the Apology of the Author, and an Essay on the Cultus of the Blessed Virgin Mary. Fourth Edition, with Facsimiles of the 5 Original Illustrations. Edited by the Rev. ORBY SHIPLEY, M.A. Fcp. 8vo. 10s. 6d.

CELEBRATED SANCTUARIES of the MADONNA. By the Rev. J. SPENCER NORTHCOTE, D.D. Post 8vo. 6s. 6d.

The HIDDEN WISDOM of CHRIST and the KEY of KNOWLEDGE; or, History of the Apocrypha. By ERNEST DE BUNSEN. 2 vols. 8vo. 28s.

The KEYS of ST. PETER; or, the House of Rechab, connected with the History of Symbolism and Idolatry. By the same Author. 8vo. 14s.

- The TYPES of GENESIS**, briefly considered as Revealing the Development of Human Nature. By ANDREW JUKES. Second Edition. Crown 8vo. 7s. 6d.
- The Second Death and the Restitution of All Things**, with some Preliminary Remarks on the Nature and Inspiration of Holy Scripture. By the same Author. Second Edition. Crown 8vo. 3s. 6d.
- ESSAYS and REVIEWS**. By the Rev. W. TEMPLE, D.D. the Rev. R. WILLIAMS, B.D. the Rev. B. POWELL, M.A. the Rev. H. B. WILSON, B.D. C. W. GOODWIN, M.A. the Rev. M. PATTISON, B.D. and the Rev. B. JOWETT, M.A. Twelfth Edition. Fcp. 8vo. 8s.
- The POWER of the SOUL over the BODY**. By GEORGE MOORE, M.D. M.R.C.P.L. &c. Sixth Edition. Crown 8vo. 8s. 6d.
- PASSING THOUGHTS on RELIGION**. By ELIZABETH M. SEWELL, Author of 'Amy Herbert.' New Edition. Fcp. 8vo. 5s.
- Self-Examination before Confirmation**. By the same Author. 32mo. price 1s. 6d.
- Readings for a Month Preparatory to Confirmation**, from Writers of the Early and English Church. By the same Author. Fcp. 4s.
- Readings for Every Day in Lent**, compiled from the Writings of Bishop JEREMY TAYLOR. By the same Author. Fcp. 5s.
- Preparation for the Holy Communion; the Devotions chiefly from the works of JEREMY TAYLOR**. By the same. 32mo. 3s.
- PRINCIPLES of EDUCATION** Drawn from Nature and Revelation, and applied to Female Education in the Upper Classes. By the Author of 'Amy Herbert.' 2 vols. fcp. 12s. 6d.
- The WIFE'S MANUAL**; or, Prayers, Thoughts, and Songs on Several Occasions of a Matron's Life. By the Rev. W. CALVERT, M.A. Crown 8vo. price 10s. 6d.
- SINGERS and SONGS of the CHURCH**: being Biographical Sketches of the Hymn-Writers in all the principal Collections; with Notes on their Psalms and Hymns. By JOSIAH MILLER, M.A. New Edition, enlarged. Crown 8vo. [Nearly ready.]
- 'SPIRITUAL SONGS' for the SUNDAYS and HOLIDAYS** throughout the Year. By J. S. B. MONSELL, LL.D. Vicar of Egham and Rural Dean. Fourth Edition, Sixth Thousand. Fcp. 4s. 6d.
- The Beatitudes**: Abasement before God; Sorrow for Sin; Meekness of Spirit; Desire for Holiness; Gentleness; Purity of Heart; the Peacemakers; Sufferings for Christ. By the same. Third Edition. Fcp. 3s. 6d.
- His PRESENCE—not his MEMORY, 1855**. By the same Author, in Memory of his SON. Sixth Edition. 16mo. 1s.
- LYRA DOMESTICA**; Christian Songs for Domestic Edification. Translated from the *Psalter* and *Harp* of C. J. P. SPITTA, and from other sources, by RICHARD MASSIE. FIRST and SECOND SERIES, fcp. 4s. 6d. each.
- LYRA GERMANICA**, translated from the German by Miss C. WINKWORTH. FIRST SERIES, Hymns for the Sundays and Chief Festivals. SECOND SERIES, the Christian Life. Fcp. 3s. 6d. each SERIES.

LYRA EUCHARISTICA; Hymns and Verses on the Holy Communion, Ancient and Modern: with other Poems. Edited by the Rev. ORBY SHIPLEY, M.A. Second Edition. Fcp. 7s. 6d.

Lyra Messianica; Hymns and Verses on the Life of Christ, Ancient and Modern; with other Poems. By the same Editor. Second Edition, altered and enlarged. Fcp. 7s. 6d.

Lyra Mystica; Hymns and Verses on Sacred Subjects, Ancient and Modern. By the same Editor. Fcp. 7s. 6d.

ENDEAVOURS after the CHRISTIAN LIFE: Discourses. By JAMES MARTINEAU. Fourth and cheaper Edition, carefully revised; the Two Series complete in One Volume. Post 8vo. 7s. 6d.

WHATELY'S Introductory Lessons on the Christian Evidences. 18mo. 6d.

INTRODUCTORY LESSONS on the HISTORY of RELIGIOUS Worship; being a Sequel to the 'Lessons on Christian Evidences.' By RICHARD WHATELY, D.D. New Edition. 18mo. 2s. 6d.

BISHOP JEREMY TAYLOR'S ENTIRE WORKS. With Life by BISHOP HEBER. Revised and corrected by the Rev. C. P. EDEN, 10 vols. price 25 5s.

Travels, Voyages, &c.

SIX MONTHS in INDIA. By MARY CARPENTER. 2 vols. post 8vo. with Portrait, 18s.

NARRATIVE of the EUPHRATES EXPEDITION carried on by Order of the British Government during the years 1835, 1836, and 1837. By General F. R. CHESNEY, F.R.S. With 2 Maps, 45 Plates, and 16 Woodcuts. 8vo. 24s.

The NORTH-WEST PENINSULA of ICELAND; being the Journal of a Tour in Iceland in the Summer of 1862. By C. W. SHEPHERD, M.A. F.Z.S. With a Map and Two Illustrations. Fcp. 8vo. 7s. 6d.

PICTURES in TYROL and Elsewhere. From a Family Sketch-Book. By the Authoress of 'A Voyage en Zigzag,' &c. Second Edition. Small 4to. with numerous Illustrations, 21s.

HOW WE SPENT the SUMMER; or, a Voyage en Zigzag in Switzerland and Tyrol with some Members of the ALPINE CLUB. From the Sketch-Book of one of the Party. In oblong 4to. with 300 Illustrations, 15s.

BEATEN TRACKS; or, Pen and Pencil Sketches in Italy. By the Authoress of 'A Voyage en Zigzag.' With 42 Plates, containing about 200 Sketches from Drawings made on the Spot. 8vo. 16s.

MAP of the CHAIN of MONT BLANC, from an actual Survey in 1863—1864. By A. ADAMS-REILLY, F.R.G.S. M.A.C. Published under the Authority of the Alpine Club. In Chromolithography on extra stout drawing-paper 25in. x 17in. price 10s. or mounted on canvas in a folding case, 12s. 6d.

HISTORY of DISCOVERY in our AUSTRALASIAN COLONIES, Australia, Tasmania, and New Zealand, from the Earliest Date to the Present Day. By WILLIAM HOWITT. 2 vols. 8vo. with 3 Maps, 20s.

- The CAPITAL of the TYCOON**; a Narrative of a Three Years' Residence in Japan. By Sir RUTHERFORD ALCOCK, K.C.B. 2 vols. 8vo. with numerous Illustrations, 42s.
- The DOLOMITE MOUNTAINS**; Excursions through Tyrol, Carinthia, Carniola, and Friuli, 1861-1863. By J. GILBERT and G. C. CHURCHILL, F.R.G.S. With numerous Illustrations. Square crown 8vo. 21s.
- GUIDE to the PYRENEES**, for the use of Mountaineers. By CHARLES PACKE. 2d Edition, with Map and Illustrations. Cr. 8vo. 7s. 6d.
- The ALPINE GUIDE**. By JOHN BALL, M.R.I.A. late President of the Alpine Club. 3 vols. post 8vo. with Maps and other Illustrations:—
- Guide to the Eastern Alps**, price 10s. 6d.
- Guide to the Western Alps**, including Mont Blanc, Monte Rosa, Zermatt, &c. Price 6s. 6d.
- Guide to the Central Alps**, including all the Oberland District. 7s. 6d.
- Introduction on Alpine Travelling in General, and on the Geology of the Alps**, price 1s. Each of the Three Volumes or Parts of the *Alpine Guide* may be had with this INTRODUCTION prefixed, price 1s. extra.
- NARRATIVES of SHIPWRECKS of the ROYAL NAVY** between 1793 and 1857, compiled from Official Documents in the Admiralty by W. O. S. GILLY; with a Preface by W. S. GILLY, D.D. Third Edition. Fcp. 5s.
- TRAVELS in ABYSSINIA and the GALLA COUNTRY**; with an Account of a Mission to Ras Ali in 1848. From the MSS. of the late WALTER CHICHELE PLOWDEN, Her Britannic Majesty's Consul in Abyssinia. Edited by his Brother TREVOR CHICHELE PLOWDEN. With Two Maps. 8vo. 18s.
- MEMORIALS of LONDON and LONDON LIFE** in the 13th, 14th, and 15th Centuries; being a Series of Extracts, Local, Social, and Political, from the Archives of the City of London. A.D. 1276-1419. Selected, translated, and edited by H. T. RILEY, M.A. Royal 8vo. 21s.
- COMMENTARIES on the HISTORY, CONSTITUTION, and CHARTERED FRANCHISES of the CITY of LONDON**. By GEORGE NORTON, formerly one of the Common Pleaders of the City of London. Third Edition. 8vo. 14s.
- CURIOSITIES of LONDON**; exhibiting the most Rare and Remarkable Objects of Interest in the Metropolis; with nearly Sixty Years' Personal Recollections. By JOHN TIMBS, F.S.A. New Edition, corrected and enlarged. 8vo. with Portrait, 21s.
- The NORTHERN HEIGHTS of LONDON**; or, Historical Associations of Hampstead, Highgate, Muswell Hill, Hornsey, and Islington. By WILLIAM HOWITT. With about 40 Woodcuts. Square crown 8vo. 21s.
- VISITS to REMARKABLE PLACES**: Old Halls, Battle-Fields, and Scenes Illustrative of Striking Passages in English History and Poetry. By WILLIAM HOWITT. 2 vols. square crown 8vo. with Woodcuts, 25s.
- The RURAL LIFE of ENGLAND**. By the same Author. With Woodcuts by Bewick and Williams. Medium 8vo. 12s. 6d.
- The IRISH in AMERICA**. By JOHN FRANCIS MAGUIRE, M.P. for Cork. Post 8vo. 12s. 6d.

ROMA SOTTERRANEA; or, an Account of the Roman Catacombs, and especially of the Cemetery of St. Callixtus. Compiled from the Works of Commendatore G. B. DE ROSSI, with the consent of the Author, by the Rev. J. S. NORTHCOTE, D.D. and the Rev. W. B. BROWNLOW. With numerous Engravings on Wood, 10 Lithographs, 10 Plates in Chromolithography, and an Atlas of Plans, all executed in Rome under the Author's superintendence for this Translation. 1 vol. 8vo. [Nearly ready.]

Works of Fiction.

The WARDEN: a Novel. By ANTHONY TROLLOPE. Crown 8vo. 2s. 6d.

Barchester Towers: a Sequel to 'The Warden.' Crown 8vo. 3s. 6d.

STORIES and TALES by ELIZABETH M. SEWELL, Author of 'Amy Herbert,' uniform Edition, each Tale or Story complete in a single Volume.

AMY HERBERT, 2s. 6d.

GERTRUDE, 2s. 6d.

EARL'S DAUGHTER, 2s. 6d.

EXPERIENCE OF LIFE, 2s. 6d.

CLEVE HALL, 3s. 6d.

IVORS, 3s. 6d.

KATHARINE ASHTON, 3s. 6d.

MARGARET PERCIVAL, 5s.

LANETON PARSONAGE, 4s. 6d.

URSULA, 4s. 6d.

A Glimpse of the World. By the Author of 'Amy Herbert,' Fcp. 7s. 6d.

The Journal of a Home Life. By the same Author. Post 8vo. 9s. 6d.

After Life; a Sequel to 'The Journal of a Home Life.' Price 10s. 6d.

UNCLE PETER'S FAIRY TALE for the XIX CENTURY. Edited by E. M. SEWELL, Author of 'Amy Herbert,' &c. Fcp. 8vo. 7s. 6d.

BECKER'S GALLUS; or, Roman Scenes of the Time of Augustus: with Notes and Excursuses. New Edition. Post 8vo. 7s. 6d.

BECKER'S CHARICLES; a Tale illustrative of Private Life among the Ancient Greeks: with Notes and Excursuses. New Edition. Post 8vo. 7s. 6d.

NOVELS and TALES by G. J. WHYTE MELVILLE:—

The GLADIATORS, 5s.

DIGBY GRAND, 5s.

KATE COVENTRY, 5s.

GENERAL BOUNCE, 5s.

HOLMBY HOUSE, 5s.

GOOD FOR NOTHING, 6s.

The QUEEN'S MARIES, 6s.

The INTERPRETER, 5s.

TALES of ANCIENT GREECE. By GEORGE W. COX, M.A. late Scholar of Trin. Coll. Oxon. Being a Collective Edition of the Author's Classical Stories and Tales, complete in One Volume. Crown 8vo. 6s. 6d.

A MANUAL of MYTHOLOGY, in the form of Question and Answer. By the same Author. Fcp. 3s.

Poetry and The Drama.

THOMAS MOORE'S POETICAL WORKS, the only Editions containing the Author's last Copyright Additions:—

SHAMROCK EDITION, crown 8vo. price 3s. 6d.

RUBY EDITION, crown 8vo. with Portrait, price 6s.

PEOPLE'S EDITION, square crown 8vo. with Portrait, &c. 12s. 6d.

LIBRARY EDITION, medium 8vo. Portrait and Vignette, 14s.

CABINET EDITION, 10 vols. fcp. 8vo. price 35s.

- MOORE'S IRISH MELODIES**, Macclise's Edition, with 161 Steel Plates from Original Drawings. Super-royal 8vo. 31s. 6d.
- Miniature Edition of Moore's Irish Melodies** with Macclise's Designs (as above) reduced in Lithography. Imp. 16mo. 10s. 6d.
- MOORE'S LALLA ROOKH**. Tenniel's Edition, with 68 Wood Engravings from original Drawings and other Illustrations. Fcp. 4to. 21s.
- SOUTHEY'S POETICAL WORKS**, with the Author's last Corrections and copyright Additions. Library Edition, in 1 vol. medium 8vo. with Portrait and Vignette, 14s. or in 10 vols. fcp. 3s. 6d. each.
- LAYS of ANCIENT ROME**; with *Ivry* and the *Armada*. By the Right Hon. LORD MACAULAY. 16mo. 4s. 6d.
- Lord Macaulay's Lays of Ancient Rome**. With 90 Illustrations on Wood, from the Antique, from Drawings by G. SCHAEF. Fcp. 4to. 21s.
- Miniature Edition of Lord Macaulay's Lays of Ancient Rome**, with the Illustrations (as above) reduced in Lithography. Imp. 16mo. 10s. 6d.
- GOLDSMITH'S POETICAL WORKS**, with Wood Engravings from Designs by Members of the ETCHING CLUB. Imperial 16mo. 7s. 6d.
- MEMORIES of some CONTEMPORARY POETS**; with Selections from their Writings. By EMILY TAYLOR. Royal 18mo. 5s.
- POEMS**. By JEAN INGELOW. Thirteenth Edition. Fcp. 8vo. 5s.
- POEMS** by Jean Ingelow. With nearly 100 Illustrations by Eminent Artists, engraved on Wood by the Brothers DALZIEL. Fcp. 4to. 21s.
- A STORY of DOOM**, and other Poems. By JEAN INGELOW. Fcp. 5s.
- POETICAL WORKS of LETITIA ELIZABETH LANDON (L.E.L.)**. 2 vols. 16mo. 10s.
- BOWDLER'S FAMILY SHAKSPEARE**, cheaper Genuine Edition, complete in 1 vol. large type, with 36 Woodcut Illustrations, price 14s. or with the same ILLUSTRATIONS, in 6 pocket vols. 3s. 6d. each.
- HORATII OPERA**, Pocket Edition, with carefully corrected Text, Marginal References, and Introduction. Edited by the Rev. J. E. YONGE, M.A. Square 18mo. 4s. 6d.
- HORATII OPERA**. Library Edition, with Marginal References and English Notes. Edited by the Rev. J. E. YONGE. 8vo. 21s.
- The ÆNEID of VIRGIL** Translated into English Verse. By JOHN CONINGTON, M.A. Crown 8vo. 9s.
- ARUNDINES CAMI**, sive Musarum Cantabrigiensium Lusus canori. Collegit atque edidit H. DRURY, M.A. Editio Sexta, curavit H. J. HODGSON, M.A. Crown 8vo. 7s. 6d.
- EIGHT COMEDIES of ARISTOPHANES**, viz. the *Acharnians*, *Knights*, *Clouds*, *Wasps*, *Peace*, *Birds*, *Frogs*, and *Plutus*. Translated into Rhymed Metres by LEONARD HAMPSON RUDD, M.A. 8vo. 15s.
- PLAYTIME with the POETS**: a Selection of the best English Poetry for the use of Children. By a LADY. Revised Edition. Crown 8vo. 5s.
- The HOLY CHILD**: a Poem in Four Cantos; also an Ode to Silence, and other Poems. By STEPHEN JENNER, M.A. Fcp. 8vo. 5s.

- The ILIAD of HOMER TRANSLATED into BLANK VERSE.** By
ICHABOD CHARLES WRIGHT, M.A. 2 vols. crown 8vo. 21s.
- The ILIAD of HOMER in ENGLISH HEXAMETER VERSE.** By
J. HENRY DART, M.A. of Exeter Coll. Oxford. Square crown 8vo. 21s.
- The ODYSSEY of HOMER.** Translated into Blank Verse by G. W.
EDGINTON, Licentiate in Medicine. Dedicated by permission to Edward
Earl of Derby. VOL. I. 8vo. with Map, 10s. 6d.
- DANTE'S DIVINE COMEDY,** translated in English Terza Rima by
JOHN DAYMAN, M.A. [With the Italian Text, after *Brunetti*, interpaged.]
8vo. 21s.
- HUNTING SONGS and MISCELLANEOUS VERSES.** By R. E.
EGERTON WARBURTON. Second Edition. Fcp. 8vo. 5s.
- An OLD STORY, and other Poems.** By ELIZABETH D. CROSS.
Second Edition. Fcp. 8vo. 3s. 6d.
- The THREE FOUNTAINS,** a Fæery Epic of Euboea; with other
Verses. By the Author of 'The Afterglow.' Fcp. 8vo. 3s. 6d.
- The Afterglow; Songs and Sonnets for my Friends.** By the Author
of 'The Three Fountains.' Second Edition. Fcp. 8vo. 5s.
- The SILVER STORE** collected from Mediæval Christian and Jewish
Mines. By the Rev. SABINE BARING-GOULD, M.A. Crown 8vo. 6s.

Rural Sports, &c.

- BLAINE'S ENCYCLOPÆDIA of RURAL SPORTS;** Hunting, Shoot-
ing, Fishing, Racing, &c. With above 600 Woodcuts (30 from Designs by
JOHN LEECH). 8vo. 42s.
- Col. HAWKER'S INSTRUCTIONS to YOUNG SPORTSMEN** in all
that relates to Guns and Shooting. Revised by the Author's SON. Square
crown 8vo. with Illustrations, 18s.
- The DEAD SHOT, or Sportsman's Complete Guide; a Treatise on**
the Use of the Gun, Dog-breaking, Pigeon-shooting, &c. By MARKSMAN.
Revised Edition. Fcp. 8vo. with Plates, 5s.
- The FLY-FISHER'S ENTOMOLOGY.** By ALFRED RONALDS. With
coloured Representations of the Natural and Artificial Insect. Sixth
Edition; with 20 coloured Plates. 8vo. 14s.
- A BOOK on ANGLING;** a complete Treatise on the Art of Angling
in every branch. By FRANCIS FRANCIS. Second Edition, with Portrait
and 15 other Plates, plain and coloured. Post 8vo. 15s.
- HANDBOOK of ANGLING:** Teaching Fly-fishing, Trolling, Bottom-
fishing, Salmon-fishing; with the Natural History of River Fish, and the
best modes of Catching them. By EPHEMERA. Fcp. Woodcuts, 5s.
- WILCOCK'S SEA-FISHERMAN;** comprising the Chief Methods of
Hook and Line Fishing in the British and other Seas, a Glance at Nets,
and Remarks on Boats and Boating. Second Edition, enlarged; with 80
Woodcuts. Post 8vo. 12s. 6d.
- The CRICKET FIELD;** or, the History and the Science of the Game
of Cricket. By JAMES PYCROFT, B.A. Fourth Edition. Fcp. 5s.

HORSE and MAN. By C. S. MARCH PHILLIPPS, Author of 'Jurisprudence,' &c. Fcp. 8vo. 2s. 6d.

The HORSE'S FOOT, and HOW to KEEP IT SOUND. By W. MILES, Esq. Ninth Edition, with Illustrations. Imperial 8vo. 12s. 6d.

A Plain Treatise on Horse-Shoeing. By the same Author. Sixth Edition. Post 8vo. with Illustrations, 2s. 6d.

Stables and Stable-Fittings. By the same. Imp. 8vo. with 13 Plates, 15s.

Remarks on Horses' Teeth, addressed to Purchasers. By the same. Post 8vo. 1s. 6d.

ROBBINS'S CAVALRY CATECHISM, or Instructions on Cavalry Exercise and Field Movements, Brigade Movements, Out-post Duty, Cavalry supporting Artillery, Artillery attached to Cavalry. 12mo. 5s.

BLAINE'S VETERINARY ART; a Treatise on the Anatomy, Physiology, and Curative Treatment of the Diseases of the Horse, Neat Cattle and Sheep. Seventh Edition, revised and enlarged by C. STEEL, M.R.C.V.S.L. 8vo. with Plates and Woodcuts, 18s.

The HORSE: with a Treatise on Draught. By WILLIAM YOUATT. New Edition, revised and enlarged. 8vo. with numerous Woodcuts, 12s. 6d.

The Dog. By the same Author. 8vo. with numerous Woodcuts, 6s.

The DOG in HEALTH and DISEASE. By STONEHENGE. With 70 Wood Engravings. Square crown 8vo. 10s. 6d.

The GREYHOUND. By STONEHENGE. Revised Edition, with 24 Portraits of Greyhounds. Square crown 8vo. 10s. 6d.

The OX; his Diseases and their Treatment: with an Essay on Parturition in the Cow. By J. R. DOBSON. Crown 8vo. with Illustrations. 7s. 6d.

Commerce, Navigation, and Mercantile Affairs.

BANKING, CURRENCY, and the EXCHANGES; a Practical Treatise. By ARTHUR CRUMP. Post 8vo. 6s.

The ELEMENTS of BANKING. By HENRY DUNNING MACLEOD, M.A. Barrister-at-Law. Post 8vo. [Nearly ready.]

The THEORY and PRACTICE of BANKING. By the same Author. Second Edition, entirely remodelled. 2 vols. 8vo. 30s.

PRACTICAL GUIDE for BRITISH SHIPMASTERS to UNITED States Ports. By PIERREPONT EDWARDS. Post 8vo. 8s. 6d.

A DICTIONARY, Practical, Theoretical, and Historical, of Commerce and Commercial Navigation. By J. R. M'CULLOCH, Esq. New and thoroughly revised Edition. 8vo. price 63s. cloth, or 70s. half-bd. in russia.

The LAW of NATIONS Considered as Independent Political Communities. By Sir TRAVERS TWISS, D.C.L. 2 vols. 8vo. 30s. or separately, PART I. *Peace*, 12s. PART II. *War*, 18s.

Works of Utility and General Information.

MODERN COOKERY for PRIVATE FAMILIES, reduced to a System of Easy Practice in a Series of carefully-tested Receipts. By ELIZA ACTON. Newly revised and enlarged Edition; with 8 Plates of Figures and 150 Woodcuts. Fcp. 6s.

On FOOD and its DIGESTION; an Introduction to Dietetics. By W. BRINTON, M.D. With 48 Woodcuts. Post 8vo. 12s.

WINE, the VINE, and the CELLAR. By THOMAS G. SHAW. Second Edition, revised and enlarged, with 32 Illustrations. 8vo. 16s.

A PRACTICAL TREATISE on BREWING; with Formulæ for Public Brewers, and Instructions for Private Families. By W. BLACK. 8vo. 10s. 6d.

SHORT WHIST. By MAJOR A. A thoroughly revised Edition, with an Essay on the Theory of the Modern Scientific Game by PROF. P. Fcp. 3s. 6d.

WHIST, WHAT TO LEAD. By CAM. Fourth Edition. 32mo. 1s.

A HANDBOOK for READERS at the BRITISH MUSEUM. By THOMAS NICHOLS. Post 8vo. 6s.

The CABINET LAWYER; a Popular Digest of the Laws of England, Civil, Criminal, and Constitutional. Twenty-fourth Edition, brought down to the close of the Parliamentary Session of 1868. Fcp. 10s. 6d.

The PHILOSOPHY of HEALTH; or, an Exposition of the Physiological and Sanitary Conditions conducive to Human Longevity and Happiness. By SOUTHWOOD SMITH, M.D. Eleventh Edition, revised and enlarged; with 113 Woodcuts. 8vo. 7s. 6d.

HINTS to MOTHERS on the MANAGEMENT of their HEALTH during the Period of Pregnancy and in the Lying-in Room. By T. BULL, M.D. Fcp. 5s.

The Maternal Management of Children in Health and Disease. By the same Author. Fcp. 6s.

The LAW RELATING to BENEFIT BUILDING SOCIETIES; with Practical Observations on the Act and all the Cases decided thereon; also a Form of Rules and Forms of Mortgages. By W. TIDD PRATT, Barrister. Second Edition. Fcp. 3s. 6d.

NOTES on HOSPITALS. By FLORENCE NIGHTINGALE. Third Edition, enlarged; with 13 Plans. Post 4to. 18s.

COULTHART'S DECIMAL INTEREST TABLES at 24 Different Rates not exceeding 5 per Cent. Calculated for the use of Bankers. To which are added Commission Tables at One-Eighth and One-Fourth per Cent. 8vo. price 15s.

MAUNDER'S TREASURY of KNOWLEDGE and LIBRARY of Reference: comprising an English Dictionary and Grammar, Universal Gazetteer, Classical Dictionary, Chronology, Law Dictionary, a Synopsis of the Peerage, useful Tables, &c. Revised Edition. Fcp. 10s. 6d.

INDEX.

Acton's Modern Cookery	24	Calvert's Wife's Manual	91
Afterglow (The).....	26	Cannon's Grant's Campaign.....	2
Alcock's Residence in Japan	23	Carpenter's Six Months in India.....	23
Alloys on Formation of Christendom.....	20	Cates's Biographical Dictionary	4
Alpine Guide (The)	22	Cats' and Fables's Moral Emblems.....	16
Apjohn's Manual of the Metalloids.....	12	Changed Aspects of Unchanged Truths.....	9
Arnold's Manual of English Literature.....	7	Chesney's Euphrates Expedition	22
Arnott's Elements of Physics.....	11	———— Indian Polity	3
Arundines Cami	25	———— Waterloo Campaign	2
Autumn holidays of a Country Parson ..	9	Child's Physiological Essays	15
Avan's Treasury of Bible Knowledge.....	20	Chorale Book for England	15
Bacon's Essays, by Whately	6	Christian Schools and Scholars	10
———— Life and Letters, by Spedding.....	5	Churchman's Daily Remembrance.....	20
———— Works, edited by Spedding.....	6	Clough's Lives from Plutarch	3
Bain's Mental and Moral Science	10	Cobb's Norman Kings of England	4
———— on the Emotions and Will.....	10	Coleenso (Bishop) on Pentateuch and Book	20
———— on the Senses and Intellect.....	10	of Joshua	20
———— on the Study of Character	7	Commonplace Philosopher in Town and	8
Ball's Alpine Guide	23	Country	8
Barnard's Drawing from Nature	16	Conington's Chemical Analysis.....	14
Baydon's Rents and Tillages.....	18	———— Translation of Virgil's <i>Æneid</i>	14
Beaten Tracks	22	CONTAINES's French-English Dictionaries	8
Becker's Charicles and Gallus	24	CONYBEARE and HOWSON's Work on St. Paul	19
BENFET'S Sanskrit Dictionary	8	COOK on the Acts.....	18
Becker's Treatise on Brewing	28	COOK'S Voyages	5
Blackley's Word-Gossip	7	COPLAND's Dictionary of Practical Medicine	15
Blackley and Friedlander's German and	8	COULTHART'S Decimal Interest Tables.....	28
English Dictionary	26	Counsel and Comfort from a City Pulpit ..	9
Blaine's Rural Sports	27	Cox's (G. W.) Manual of Mythology	24
———— Veterinary Art.....	9	———— Tale of the Great Persian War ..	2
Booth's Epigrams.....	18	———— Tales of Ancient Greece	24
Bourne on Screw Propeller	17	———— (H.) Ancient Parliamentary Elections	1
Bourne's Catechism of the Steam Engine..	17	———— History of the Reform Bills	1
———— Handbook of Steam Engine.....	17	———— Whig and Tory Administrations ..	1
———— Treatise on the Steam Engine.....	17	Cressy's Encyclopedia of Civil Engineering	17
———— Examples of Modern Engines ..	13	Critical Essays of a Country Parson.....	9
Bowdler's Family SHAKESPEARE	25	Cross's Old Story	26
Bowdler's Dictionary of Science, Literature,	13	Crowe's History of France	3
and Art.....	10	Crimp on Banking, Currency, & Exchanges	27
Brat's (C.) Education of the Feelings.....	10	Croley's Handbook of Telegraphy	17
———— Philosophy of Necessity.....	10	Cusack's History of Ireland.....	3
———— on Force	10	Dart's Iliad of Homer.....	26
Brainton on Food and Digestion.....	28	D'Aubigne's History of the Reformation in	2
Brodie's (Sir C. B.) Works.....	15	the time of CALVIN.....	19
Brownie's Exposition of the 39 Articles ..	18	Davidson's Introduction to New Testament	2
Buckle's History of Civilization	2	Dayman's Dante's Divine Commedia	28
Bull's Hints to Mothers.....	26	Dead Shot (The), by MARRISMAN	26
———— Maternal Management of Children.	28	De la Rive's Treatise on Electricity	11
Bunsen's (Baron) Ancient Egypt	3	De Morgan on Matter and Spirit	9
———— God in History	3	De Tocqueville's Democracy in America..	2
———— Memoirs	4	Dorson on the Ox	27
Bunsen (E. De) on Apocrypha.....	20	Dove on Storms	11
———— 's Keys of St. Peter	20	Dyer's City of Rome	3
Burke's Vicissitudes of Families	5	Eastlake's Hints on Household Taste	17
Burton's Christian Church	4	———— History of Oil Painting	15
Cabinet Lawyer	28		

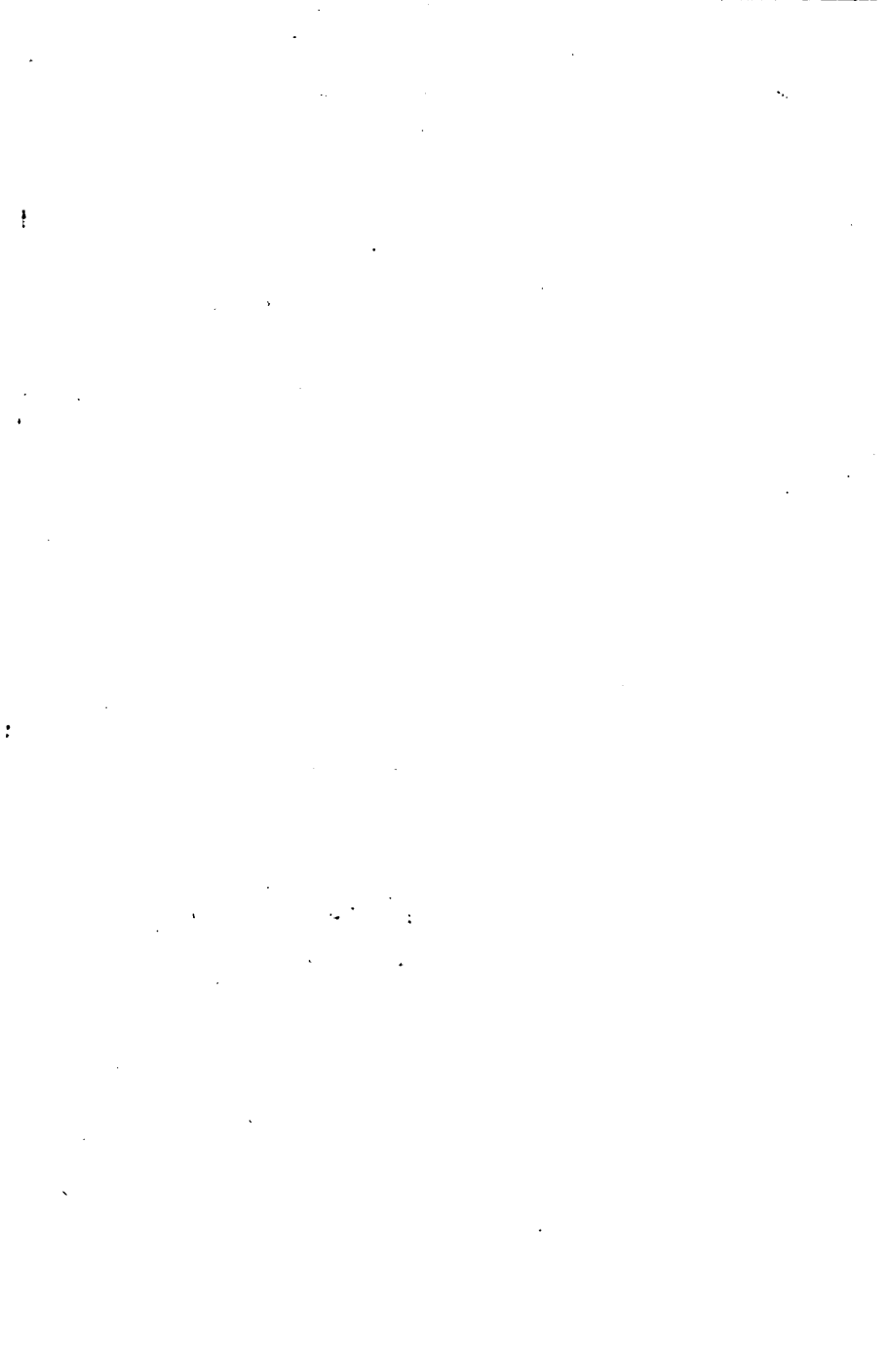
EDMUNTON'S <i>Odyssey</i>	26	HULLAR'S <i>Transition Musical Lectures</i>	15
EDWARDS'S <i>Shipmaster's Guide</i>	27	HUMPHREYS' <i>Sentiments of Shakspeare</i>	16
ELEMENTS OF Botany	18		
ELLICOTT'S <i>Commentary on Ephesians</i>	19		
<i>Lectures on Life of Christ</i>	19	INGLELOW'S <i>Poems</i>	25
<i>Commentary on Galatians</i>	19	<i>Story of Doom</i>	25
<i>Pastoral Epist.</i>	19		
<i>Philippians, &c.</i>	19	JAMESON'S <i>Legends of the Saints and Mar-</i>	
<i>Thessalonians</i>	19	<i>tyrs</i>	16
ESSAYS AND CONTRIBUTIONS OF A. K. H. B. . .	8	<i>Legends of the Madonna</i>	16
ESSAYS AND REVIEWS	21	<i>Legends of the Monastic Orders</i>	16
EWALD'S <i>History of Israel</i>	19	JAMESON AND EASTLAKE'S <i>History of Our</i>	
		<i>Lord</i>	16
FAIRBAIRN ON <i>Iron Shipbuilding</i>	17	JENNER'S <i>Holy Child</i>	25
FAIRBAIRN'S <i>Application of Cast and</i>		JOHNSTON'S <i>Gazetteer, or Geographical Dic-</i>	
<i>Wrought Iron to Building</i>	17	<i>tionary</i>	11
<i>Information for Engineers</i>	17	JORDAN ON <i>Via Inertia in the Ocean</i>	11
<i>Treatise on Mills & Millwork</i>	17	JUKES ON <i>Second Death</i>	21
FARRAR'S <i>Chapters on Language</i>	7	<i>on Types of Genesis</i>	21
FELKIN ON <i>Hosiery and Lace Manufactures</i>	18		
FIDDLER'S <i>Christendom's Divisions</i>	10	KALISCH'S <i>Commentary on the Bible</i>	7
FITZGERALD'S <i>Ireland</i>	3	<i>Hebrew Grammar</i>	7
FLIEDNER'S (<i>Pastor</i>) <i>Life</i>	5	KEITH ON <i>Fulfillment of Prophecy</i>	19
FORBES'S <i>Earls of Granard</i>	5	<i>Destiny of the World</i>	19
FRANCIS'S <i>Fishing Book</i>	26	KERL'S <i>Metallurgy by CROOKES and RÖHMIG</i>	18
FRONDE'S <i>History of England</i>	1	KESSEY'S <i>Domestic Medicine</i>	15
<i>Short Studies on Great Subjects</i>	9	KIRBY AND SPENCER'S <i>Entomology</i>	13
GANOT'S <i>Elementary Physics</i>	11	LANDON'S (<i>L. E. L.</i>) <i>Poetical Works</i>	25
GILBERT AND CHURCHILL'S <i>Dolomite Moun-</i>		LATHAM'S <i>English Dictionary</i>	7
<i>tains</i>	23	<i>River Plate</i>	11
GILLY'S <i>Shipwrecks of the Navy</i>	23	LUCKY'S <i>History of European Morals</i>	3
GOLDMITS'S <i>Poems, Illustrated</i>	25	<i>Rationalism</i>	3
GOODWIN'S <i>Elements of Mechanism</i>	17	LEIGHTON'S <i>Sermons and Charges</i>	18
GOULD'S <i>Silver Store</i>	26	<i>Leisure Hours in Town</i>	8
GRAHAM'S <i>Book about Words</i>	7	LESSONS OF <i>Middle Age</i>	9
GRANT'S <i>Ethics of Aristotle</i>	6	LEWIS' <i>History of Philosophy</i>	3
GRATER <i>Thoughts of a Country Parson</i>	9	LIDDELL AND SCOTT'S <i>Greek-English Lexicon</i>	8
GRAY'S <i>Anatomy</i>	14	<i>Abridged ditto</i>	8
GREENE'S <i>Coals and Sea Jellies</i>	12	<i>Life of Man Symbolised</i>	16
<i>Sponges and Animalcules</i>	12	LINDLEY AND MOORE'S <i>Treasury of Botany</i>	13
GREENHOW ON <i>Bronchitis</i>	14	LONGMAN'S <i>Edward the Third</i>	2
GROVE ON <i>Correlation of Physical Forces</i>	12	<i>Lectures on the History of Eng-</i>	
GWYN'S <i>Encyclopedia of Architecture</i>	16	<i>land</i>	2
		LOUDON'S <i>Agriculture</i>	18
HANDBOOK OF <i>Angling</i> , by EFFENBERG	26	<i>Gardening</i>	19
HARRIS ON <i>Election of Representatives</i>	7	<i>Plants</i>	13
HARTWIG'S <i>Harmonies of Nature</i>	13	LOWNDES'S <i>Engineer's Handbook</i>	17
<i>Polar World</i>	13	LYRA <i>Domestica</i>	21
<i>Sea and its Living Wonders</i>	13	<i>Eucharistica</i>	22
<i>Tropical World</i>	13	<i>Germanica</i>	16, 22
HAUGHTON'S <i>Manual of Geology</i>	12	<i>Messianica</i>	22
HAWKES'S <i>Instructions to Young Sportsmen</i>	26	<i>Mystica</i>	22
HENDERSON'S <i>Folk-Lore of the Northern</i>			
<i>Counties</i>	10	MACAULAY'S (<i>Lord</i>) <i>Essays</i>	3
HERSCHEL'S <i>Outlines of Astronomy</i>	10	<i>History of England</i>	1
HEWITT ON <i>Diseases of Women</i>	14	<i>Lays of Ancient Rome</i>	25
HODGSON'S <i>Time and Space</i>	9	<i>Miscellaneous Writings</i>	9
HOLMES'S <i>System of Surgery</i>	14	<i>Speeches</i>	7
<i>Surgical Diseases of Infancy</i>	14	<i>Complete Works</i>	1
HOOKE AND WALKER-ARNOTT'S <i>British</i>		MACFARREN'S <i>Lectures on Harmony</i>	15
<i>Flora</i>	13	MACLEOD'S <i>Elements of Political Economy</i>	7
HOBBS'S <i>Introduction to the Scriptures</i>	19	<i>Dictionary of Political Economy</i>	7
<i>Compendium of ditto</i>	19	<i>Elements of Banking</i>	27
HORSLEY'S <i>Manual of Poisons</i>	15	<i>Theory and Practice of Banking</i>	27
HOW WE SPENT THE <i>Summer</i>	22	McCULLOCH'S <i>Dictionary of Commerce</i>	27
HOWARD'S <i>Gymnastic Exercises</i>	15	<i>Geographical Dictionary</i>	11
HOWITT'S <i>Australian Discovery</i>	22	MAQUIR'S <i>Irish in America</i>	23
<i>Northern Heights of London</i>	23	<i>Life of Father Mathew</i>	4
<i>Rural Life of England</i>	23	MALLETON'S <i>French in India</i>	3
<i>Visits to Remarkable Places</i>	23	MANNING'S <i>England and Christendom</i>	20
HUGHES'S (<i>W.</i>) <i>Manual of Geography</i>	11	MARSHALL'S <i>Physiology</i>	14
HULLAR'S <i>Collection of Sacred Music</i>	16	MARSHMAN'S <i>Life of Havelock</i>	5
<i>Lectures on Modern Music</i>	15	<i>History of India</i>	3

MARTINEAU'S Endeavours after the Christian Life	22	ODLING'S Manual of Chemistry	14
MASSEY'S History of England	2	Original Designs for Wood Carving	17
MASSINGBERD'S History of the Reformation ..	4	OWEN'S Lectures on the Invertebrate Animals	12
MAUNDER'S Biographical Treasury	5	Comparative Anatomy and Physiology of Vertebrate Animals	13
Geographical Treasury	11		
Historical Treasury	13	PAGET'S Guide to the Pyrenees	23
Scientific and Literary Treasury	28	PAGET'S Lectures on Surgical Pathology	14
Treasury of Knowledge	13	PERRIRA'S Manual of Materia Medica	23
Treasury of Natural History	13	PERRINS'S Italian and Tuscan Sculptors	17
MAURY'S Physical Geography	11	PHILLIPS'S Guide to Geology	13
MAT'S Constitutional History of England ..	2	PHILLIPS'S Horse and Man	27
MEISNER'S Biographical and Critical Essays ..	5	Pictures in Tyrol	23
MELIA ON Virgin Mary	20	PISSAS'S Art of Perfumery	18
MELVILLE'S Digby Grand	24	Chemical, Natural, and Physical Magic	18
General Bounce	24	PIKE'S English and their Origin	9
Gladiators	21	Playtime with the Poets	25
Good for Nothing	24	PLowden'S Travels in Abyssinia	23
Holmby House	24	POLKO'S Reminiscences of Mendelssohn	5
Interpreter	24	PRATT'S Law of Building Societies	28
Kate Coventry	24	PRESCOTT'S Scripture Difficulties	20
Queen's Maries	24	PROCTOR'S Saturn	16
MENDLSOHN'S Letters	5	Handbook of the Stars	10
MERIVALE'S (H.) Historical Studies	2	PROCTOR'S Cricket Field	28
(C.) Fall of the Roman Republic	3		
Romans under the Empire	10	Quarterly Journal of Science	13
MERRIFIELD AND EVER'S Navigation	27	QUICK'S Educational Reformers	5
MILES ON Horse's Foot and Horseshoeing ..	27		
MILES ON Horse's Teeth and Stables	27	Recreations of a Country Parson	8
MILL (J.) on the Mind	10	RILEY'S Map of Mont Blanc	23
MILL (J. S.) on Liberty	6	REIMANN ON Aniline Dyes	15
on Representative Government	6	RICHARDSON'S Life, by McILWRAITH	5
on Utilitarianism	6	RILEY'S Memorials of London	23
MILL'S (J. S.) Dissertations and Discussions ..	6	RIVERS'S Rose Amateur's Guide	13
Political Economy	6	ROBBINS'S Cavalry Catechism	27
System of Logic	6	ROGERS'S Correspondence of Greyson	9
Hamilton's Philosophy	7	Eclipse of Faith	9
Inaugural Address	7	Defence of ditto	9
MILLER'S Elements of Chemistry	14	Essays from the <i>Edinburgh Review</i>	9
Hymn-Writers	21	Reason and Faith	9
MITCHELL'S Manual of Assaying	18	ROBERT'S Thesaurus of English Words and Phrases	7
Modern Ireland	3	Roma Sotteranea	24
MONKELL'S Beatitudes	21	RONALD'S Fly-Fisher's Entomology	26
His Presence not his Memory	21	ROWTON'S Debater	7
Spiritual Songs	21	RUON'S Aristophanes	25
MOORE'S Irish Melodies	25	RUSSELL ON Government and Constitution ..	1
Lalla Rookh	24		
Poetical Works	24	SANDARS'S Justinian's Institutes	6
Dr. G.'s First Man	12	SCHAEFFLER ON Ocular Defects and Spectacles ..	14
Power of the Soul over the Body ..	21	SCHERRER'S Life, translated by COLERIDGE ..	4
MORRELL'S Elements of Psychology	10	SCOTT'S Lectures on the Fine Arts	15
Mental Philosophy	10	SEABORN'S Oxford Reformers of 1488	2
MOUNTFIELD ON National Church	19	SEWELL'S Journals &c. relating to Ireland ..	3
MÜLLER'S (MAX) Chips from a German Workshop	9	SEWELL'S After Life	24
Lectures on the Science of Language	7	Amy Herbert	24
(K. O.) Literature of Ancient Greece	2	Cleve Hall	24
MURCHISON ON Continued Fevers	14	Earl's Daughter	24
on Liver Complaints	14	Examination for Confirmation	21
MURRAY'S Language and Literature of Greece ..	3	Experience of Life	24
		Gertrude	24
New Testament, Illustrated with Wood Engravings from the Old Masters	16	Glimpse of the World	4
NEWMAN'S History of his Religious Opinions ..	4	History of the Early Church	4
NICHOLAS'S Pedigree of the English People ..	28	Ivory	24
NICHOLS'S Handbook to the British Museum ..	28	Journal of a Home Life	24
NORTHING'S Notes on Hospitals	23	Katharine Ashton	24
NILSON'S Scandinavia	12	Janetson Parsonage	24
NORTHCOOTE'S Sanctuaries of the Madonna ..	30	Margaret Percival	24
NORTHCOOTE'S Lathes and Turning	17	Passing Thoughts on Religion	21
NORTON'S City of London	23	Preparation for Communion	21
		Principles of Education	21
ODLING'S Animal Chemistry	14	Readings for Confirmation	21
Course of Practical Chemistry	14	Readings for Lent	21
		Tales and Stories	23

SEWELL'S Ursula	24	TINDALL'S Lectures on Heat.....	11
SEAW'S Work on Wine	28	Sound	11
SHAKESPEARE'S Midsummer Night's Dream illustrated with Silhouettes	16	FARADAY AS A Discoverer	5
SHEPHERD'S Iceland	22	UNCLE PETER'S Fairy Tale	24
SHIPLEY'S Church and the World	20	URE'S Dictionary of Arts, Manufactures, and Mines	17
SHORT Whist	28	VAN DER HOEVEN'S Handbook of Zoology..	12
SHORT'S Church History	4	WARBURTON'S Hunting Songs	26
SMART'S WALKER'S Pronouncing Dictionary ..	2	WATSON'S Principles and Practice of Physic ..	14
SMITH'S (SOUTHWOOD) Philosophy of Health ..	8	WATTS'S Dictionary of Chemistry	13
(J.) Paul's Voyage and Shipwreck..	19	WEBB'S Objects for Common Telescopes....	10
(SYDNEY) Miscellaneous Works	9	WEBSTER & WILKINSON'S Greek Testament ..	20
Wit and Wisdom	9	WELD'S Florence	22
SOUTHEY'S (Doctor)	7	WELLINGTON'S Life, by the Rev G. R. GLEIG	4
Poetical Works	25	WELLS ON DEW	11
STAFFORD'S Life of the Blessed Virgin	20	WEST ON Children's Diseases	14
STANLEY'S History of British Birds	12	WHATELY'S English Synonyms	6
STEEBING'S Analysis of MILL'S Logic	6	Logic	6
STEPHEN'S Essays in Ecclesiastical Bio- graphy	5	Rhetoric	6
STIRLING'S Secret of Hegel	10	Life and Correspondence	4
STORES'S Life of PETRÆ	5	WHATELY on the Truth of Christianity....	22
STONEHENGE on the Dog	27	Religious Worship	22
on the Greyhound	27	Whist, what to lead, by CAM	28
STRICKLAND'S Tudor Princesses	4	WHITE and RIDDLE'S Latin-English Dic- tionaries	8
Sunday Afternoons at the Parish Church of a Scottish University City (St. Andrew's) ..	9	WILCOCKS'S Sea Fisherman	26
TAYLOR'S (Jeremy) Works, edited by EDEN ..	22	WINSLOW ON Light	11
(E.) Selections from some Contem- porary Poets	25	WOOD'S Bible Animals	12
TENNENT'S Ceylon	13	Homes without Hands	12
THIRLWALL'S History of Greece	2	WRIGHT'S Homer's Iliad	26
THOMSON'S (Archbishop) Laws of Thought ..	6	YEO'S Manual of Zoology	12
(A. T.) Conspectus	15	YONGE'S English-Greek Lexicons	9
Three Fountains (The)	26	Editions of Horace	25
TIMBS'S Curiosities of London	23	YOCATT on the Dog	27
TODD (A.) on Parliamentary Government..	1	on the Horse	27
TODD and BOWMAN'S Anatomy and Phy- siology of Man	15	ZELLER'S Socrates	6
TRENCH'S Realities of Irish Life	3		
TROLLOPE'S Barchester Towers	24		
Warden	24		
Twiss's Law of Nations	27		



LONDON: PRINTED BY
SPOTTISWOODE AND CO., NEW-STREET SQUARE
AND PARLIAMENT STREET



14 DAY USE

RETURN TO DESK FROM WHICH BORROWED

LOAN DEPT.

**This book is due on the last date stamped below, or
on the date to which renewed.**

Renewed books are subject to immediate recall.

ICLF (N)

LD 21A-60m-2, '67
(H241s10)476B

General Library
University of California
Berkeley

YB 17072

LIBRARY USE
RETURN TO DESK FROM WHICH BORROWED

LOAN DEPT.

THIS BOOK IS DUE BEFORE CLOSING TIME
ON LAST DATE STAMPED BELOW

ICLF (N)

LIBRARY USE

JAN 24 1968

REC'D LD

JAN 24 '68 - 9 AM

General Library
University of California
Berkeley

